



# **STIC Search Report**

## **Biotech-Chem Library**

**STIC Database Tracking Number: 118581**

**To: Terra Gibbs**  
**Location: REM/2D10/2C18**  
**Art Unit: 1635**  
**Wednesday, April 07, 2004**

**Case Serial Number: 10/033243**

**From: Beverly Shears**  
**Location: Remsen Bldg.**  
**RM 1A54**  
**Phone: 571-272-2528**

**[beverly.shears@uspto.gov](mailto:beverly.shears@uspto.gov)**

### **Search Notes**

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STIC-Biotech/ChemLib

118581

From: Gibbs, Terra  
Sent: Friday, April 02, 2004 2:33 PM  
To: STIC-Biotech/ChemLib  
Subject: Sequence search...

Can you please do a search of SEQ ID NO:132 of USSN 10/033,243 in all regular and commercial databases?

Terra Cotta Gibbs, Ph.D.  
Art Unit 1635  
Remsen Building 2D10  
571-272-0758

RECEIVED  
APR -2 2004  
STIC-Biotech/ChemLib  
(STIC)

Searcher: \_\_\_\_\_  
Phone: \_\_\_\_\_  
Location: \_\_\_\_\_  
Date Picked Up: \_\_\_\_\_  
Date Completed: \_\_\_\_\_  
Searcher Prep/Review: \_\_\_\_\_  
Clerical: \_\_\_\_\_  
Online time: \_\_\_\_\_

TYPE OF SEARCH:  
NA Sequences: \_\_\_\_\_  
AA Sequences: \_\_\_\_\_  
Structures: \_\_\_\_\_  
Bibliographic: \_\_\_\_\_  
Litigation: \_\_\_\_\_  
Full text: \_\_\_\_\_  
Patent Family: \_\_\_\_\_  
Other: \_\_\_\_\_

VENDOR/COST (where applic.)  
STN: \_\_\_\_\_  
DIALOG: \_\_\_\_\_  
Questel/Orbit: \_\_\_\_\_  
DRLink: \_\_\_\_\_  
Lexis/Nexis: \_\_\_\_\_  
Sequence Sys.: \_\_\_\_\_  
WWW/Internet: \_\_\_\_\_  
Other (specify): \_\_\_\_\_

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# SEARCH REQUEST FORM

Requestor's Name: \_\_\_\_\_ Serial Number: \_\_\_\_\_

Date: \_\_\_\_\_ Phone: \_\_\_\_\_ Art Unit: \_\_\_\_\_

## Search Topic:

Please write a detailed statement of search topic. Describe specifically as possible the subject matter to be searched. Define any terms that may have a special meaning. Give examples or relevant citations, authors, keywords, etc., if known. For sequences, please attach a copy of the sequence. You may include a copy of the broadest and/or most relevant claim(s).

## STAFF USE ONLY

Date completed: 04-06-04  
 Searcher: Beverly C 2528  
 Terminal time: 22  
 Elapsed time: \_\_\_\_\_  
 CPU time: \_\_\_\_\_  
 Total time: 25  
 Number of Searches: \_\_\_\_\_  
 Number of Databases: 2

### Search Site

\_\_\_\_\_ STIC  
 \_\_\_\_\_ CM-1  
 \_\_\_\_\_ Pre-S

### Type of Search

\_\_\_\_\_ N.A. Sequence  
 \_\_\_\_\_ A.A. Sequence  
 \_\_\_\_\_ Structure  
 \_\_\_\_\_ Bibliographic

### Vendors

\_\_\_\_\_ IG  
☒ STN  
 \_\_\_\_\_ Dialog  
 \_\_\_\_\_ APS  
 \_\_\_\_\_ Geninfo  
 \_\_\_\_\_ SDC  
 \_\_\_\_\_ DARC/Questel  
☒ Other CGN

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10/033243

L1 FILE 'REGISTRY' ENTERED AT 08:18:30 ON 06 APR 2004  
26 S TCGTCGAACGTTTCGAGATGAT/SQSN

L2 FILE 'HCAPLUS' ENTERED AT 08:19:36 ON 06 APR 2004  
7 S L1

L2 ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2004 ACS on STN  
ED Entered STN: 22 Feb 2004  
ACCESSION NUMBER: 2004:142919 HCAPLUS  
DOCUMENT NUMBER: 140:198064  
TITLE: Particulate immunostimulant  
INVENTOR(S): Van Nest, Gary; Tuck, Stephen  
PATENT ASSIGNEE(S): Dynavax Technologies Corporation, USA  
SOURCE: PCT Int. Appl., 90 pp.  
CODEN: PIXXD2  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004014322	A2	20040219	WO 2003-US25415	20030812
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

PRIORITY APPLN. INFO.: US 2002-402968P P 20020812  
AB The authors disclose immunomodulatory compns. which comprise a cationic condensing agent, an immunomodulatory compound, and a stabilizing agent. The compns. of the invention typically form particles which have increased immunomodulatory activity as compared to immunomodulatory compds. not formulated in the compns. of the invention. Also provided are methods of making the compns. and methods for therapeutic use of the compns. In one example, interferon- $\gamma$  release by human mononuclear cells was shown to be enhanced by the combination of CpG oligonucleotide, polymyxin B, and Tween-80.

IT 662376-82-1  
RL: PRP (Properties)  
(unclaimed sequence; particulate immunostimulant)

L2 ANSWER 2 OF 7 HCAPLUS COPYRIGHT 2004 ACS on STN  
ED Entered STN: 15 Dec 2003  
ACCESSION NUMBER: 2003:975836 HCAPLUS  
DOCUMENT NUMBER: 140:75912  
TITLE: IL-10 regulates plasmacytoid dendritic cell response to CpG-containing immunostimulatory sequences

Searcher : Shears 571-272-2528

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10/033243

AUTHOR(S): Duramad, Omar; Fearon, Karen L.; Chan, Jean H.;  
Kanzler, Holger; Marshall, Jason D.; Coffman,  
Robert L.; Barrat, Franck J.

CORPORATE SOURCE: Dynavax Technologies Corporation, Berkeley, CA,  
USA

SOURCE: Blood (2003), 102(13), 4487-4492  
CODEN: BLOOAW; ISSN: 0006-4971

PUBLISHER: American Society of Hematology

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Immunostimulatory sequences (ISS) are short oligonucleotides containing unmethylated cytosine-phosphate-guanine (CpG) dinucleotides that stimulate innate immune responses through Toll-like receptor-9 on B cells and plasmacytoid dendritic cell (PDC) precursors. The anti-inflammatory cytokine interleukin (IL)-10 is predicted to be a potent inhibitor of many of the activities described for ISS, and this may impact the use of ISS in disease states characterized by elevated IL-10. As the activities of ISS on PDCs are central to many clin. applications of ISS, we have studied the effects of IL-10 on PDC stimulation by 3 distinct classes of ISS. IL-10 inhibited cytokine production and survival of ISS-activated PDCs; however, IL-12 induction was much more sensitive to inhibition than interferon (IFN)- $\alpha$  induction. Within the PDC population are cells that respond to ISS by producing either IL-12 or IFN- $\alpha$  but not both cytokines. IL-12-producing PDCs require costimulation through CD40 and appear more mature than IFN- $\alpha$ -producing PDCs. The 3 distinct classes of ISS differed with respect to induction of PDC maturation and T-cell priming capacity. IL-10 regulated PDC activation but did not inhibit the subsequent T-cell-priming ability of PDCs already activated by ISS.

IT 640803-45-8  
RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); BIOL (Biological study)  
(IL-10 regulates plasmacytoid dendritic cell response to CpG-containing immunostimulatory sequences)

REFERENCE COUNT: 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 3 OF 7 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 05 Dec 2003

ACCESSION NUMBER: 2003:950038 HCAPLUS

DOCUMENT NUMBER: 140:26897

TITLE: Chimeric immunomodulatory compounds comprising two or more nucleic acid moieties and non-nucleic acid spacer

INVENTOR(S): Fearon, Karen L.; Dina, Dino; Tuck, Stephen F.

PATENT ASSIGNEE(S): USA

SOURCE: U.S. Pat. Appl. Publ., 96 pp., Cont.-in-part of U.S. Ser. No. 176,883.  
CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

Searcher : Shears 571-272-2528

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PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003225016	A1	20031204	US 2002-328578	20021223
US 2003175731	A1	20030918	US 2002-176883	20020621
US 2003199466	A1	20031023	US 2002-177826	20020621

PRIORITY APPLN. INFO.:

US 2001-299883P	P	20010621
US 2002-375253P	P	20020423
US 2002-176883	A2	20020621
US 2002-177826	A2	20020621

AB The invention provides immunomodulatory compds. and methods for immunomodulation of individuals using the immunomodulatory compds. The immunomodulatory compds. comprise two or more nucleic acid moieties and a non-nucleic acid spacer moiety. The nucleic acid contains e.g. 5'-CG-3', 5'-TCG-3', 5'-TCGA-3', 5'-TCGACGT-3', or 5'-TCGACGA-3'; and the non-nucleic acid is an oligoethylene glycol such as hexaethylene glycol. The chimeric compds. are incorporated into endotoxin-free compns. comprising antigen, pharmaceutically acceptable excipient, and optionally a cationic microsphere for modulating immune response.

IT 631925-51-4P 631925-64-9P 631926-21-1P  
631926-36-8P  
RL: PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
(chimeric immunomodulatory compds. comprising two or more nucleic acid moieties and non-nucleic acid spacer)

L2 ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2004 ACS on STN  
ED Entered STN: 01 Jul 2003  
ACCESSION NUMBER: 2003:499872 HCAPLUS  
DOCUMENT NUMBER: 139:99590  
TITLE: Rational design of new CpG oligonucleotides that combine B cell activation with high IFN- $\alpha$  induction in plasmacytoid dendritic cells  
AUTHOR(S): Hartmann, Gunther; Battiany, Julia; Poeck, Hendrik; Wagner, Moritz; Kerkmann, Miren; Lubenow, Norbert; Rothenfusser, Simon; Endres, Stefan  
CORPORATE SOURCE: Department of Internal Medicine, Division of Clinical Pharmacology, Ludwig-Maximilians-University of Munich, Munich, Germany  
SOURCE: European Journal of Immunology (2003), 33(6), 1633-1641  
CODEN: EJIMAF; ISSN: 0014-2980  
PUBLISHER: Wiley-VCH Verlag GmbH & Co. KGaA  
DOCUMENT TYPE: Journal  
LANGUAGE: English

AB Two different types of CpG motif-containing oligonucleotides (CpG ODN) have been described: CpG-A with high induction of IFN- $\alpha$  in plasmacytoid dendritic cells; and CpG-B with little induction of IFN- $\alpha$ , but potent activation of B cells. In this study, we demonstrate that CpG-A fail to activate B cells unless plasmacytoid dendritic cells are present. We identified a new set of CpG ODN sequences which induces high levels of IFN- $\alpha$  in plasmacytoid dendritic cells but remains capable of directly activating B cells. These new CpG ODN (termed CpG-C) are more potent stimulants of B cells than CpG-B due to their ability of directly and indirectly

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(via plasmacytoid dendritic cells) activating B cells. The sequence of CpG-C combines structural elements of both CpG-A and CpG-B. The most potent sequence, M362, contains a 5'-end "TCGTCG-motif" and a "GTCGTT-motif", both of which are present in CpG-B (ODN 2006); a palindromic sequence characteristic for CpG-A (ODN 2216); but no poly G motif required for CpG-A. In conclusion, we defined the first CpG-containing sequences that potentially activate both TLR9-expressing immune cell subsets in humans, the plasmacytoid dendritic cell and the B cell. CpG-C may allow for improved therapeutic immuno-modulation in vivo.

IT 557124-76-2P

RL: BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)  
(rational design of new CpG oligonucleotides that combine B cell activation with high IFN- $\alpha$  induction in plasmacytoid dendritic cells)

REFERENCE COUNT: 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 13 Jun 2003

ACCESSION NUMBER: 2003:453345 HCAPLUS

DOCUMENT NUMBER: 139:83598

TITLE: Identification of a novel CpG DNA class and motif that optimally stimulate B cell and plasmacytoid dendritic cell functions

AUTHOR(S): Marshall, Jason D.; Fearon, Karen; Abbate, Christi; Subramanian, Sandhya; Yee, Priscilla; Gregorio, Josh; Coffman, Robert L.; Van Nest, Gary

CORPORATE SOURCE: Dynavax Technologies Corporation, Berkeley, CA, USA

SOURCE: Journal of Leukocyte Biology (2003), 73(6), 781-792

CODEN: JLBIE7; ISSN: 0741-5400

PUBLISHER: Federation of American Societies for Experimental Biology

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Recent reports have identified two major classes of CpG motif-containing oligodeoxynucleotide immunostimulatory sequences (ISS): uniformly modified phosphorothioate (PS) oligodeoxyribonucleotides (ODNs), which initiate B cell functions but poorly activate dendritic cells (DCs) to make interferon (IFN)- $\alpha$ , and chimeric PS/phosphodiester (PO) ODNs containing runs of six contiguous guanosines, which induce very high levels of plasmacytoid DC (PDC)-derived IFN- $\alpha$  but poorly stimulate B cells. The authors have generated the first reported ISS, C274, which exhibits very potent effects on all human immune cells known to recognize ISS. C274 is a potent inducer of IFN- $\gamma$ /IFN- $\alpha$  from peripheral blood mononuclear cells and exhibits accelerated kinetics of activity compared with standard ISS. This ODN also effectively stimulates B cells to proliferate, secrete cytokines, and express costimulatory antigens. In addition, C274 specifically activates PDCs to undergo maturation and secrete cytokines, including very high levels of IFN- $\alpha$ . Sequence

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variation studies based on C274 were used to identify the general motif requirements for this novel and distinct class of ISS. In contrast, chimeric PO/PS CpG-containing ODNs with Poly guanosine sequences exert a differential pattern of ISS activity compared with C274, perhaps in part as a result of their greatly different structural nature. This pattern is composed of high IFN- $\alpha$ /IFN- $\gamma$  induction and low DC maturation in the absence of B cell stimulation. In conclusion, the authors have generated a novel class of ISS that transcends the limitations ascribed to classes described previously in that it provides excellent stimulation of B cells and simultaneously activates PDCs to differentiate and secrete large amts. of type I IFN.

IT 554470-62-1P

RL: BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)  
(identification of a novel CpG DNA class and motif that optimally stimulate B cell and plasmacytoid dendritic cell functions)

REFERENCE COUNT: 61 THERE ARE 61 CITED REFERENCES AVAILABLE  
FOR THIS RECORD. ALL CITATIONS AVAILABLE  
IN THE RE FORMAT

L2 ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 05 Jan 2003

ACCESSION NUMBER: 2003:6160 HCAPLUS

DOCUMENT NUMBER: 138:88635

TITLE: Chimeric immunomodulatory compounds comprising  
nucleic acids linked through dendrimer or  
polysaccharide spacer and antigen for treating  
allergy, infection or cancer

INVENTOR(S): Fearon, Karen L.; Dina, Dino; Tuck, Stephen F.

PATENT ASSIGNEE(S): Dynavax Technologies Corporation, USA

SOURCE: PCT Int. Appl., 224 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003000922	A2	20030103	WO 2002-US20025	20020621
WO 2003000922	A3	20031023		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

PRIORITY APPLN. INFO.: US 2001-299883P P 20010621

US 2002-375253P P 20020423

AB The invention provides immunomodulatory compds. (CIC) and methods

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for immunomodulation of individuals using the immunomodulatory compds. The CIC comprises one or more nucleic acid moieties and one or more non-nucleic acid moieties such as dendrimer, polysaccharide, and crosslinked polysaccharide through phosphodiester, phosphorothioate ester, phosphorodithioate ester, and other linkages. The CIC is capable of stimulating production of interferon  $\gamma$  and  $\alpha$  by human peripheral blood mononuclear cells, as well as human B cell proliferation. Endotoxin-free compns. comprising the CIC covalently or non-covalently conjugated with antigen and cationic microsphere are useful for treating disorders associated with IgE or Th2-type immune response such as allergy, asthma, infection, viral infection, idiopathic pulmonary fibrosis, and cancer.

IT 482661-47-2P 482663-36-5P 482663-37-6P  
482663-46-7P 482663-95-6P 482663-96-7P  
482663-97-8P 483382-56-5P 483382-57-6P  
483382-58-7P 483382-59-8P 483382-64-5P  
483382-65-6P 483382-66-7P 483382-67-8P  
483382-68-9P

RL: PAC (Pharmacological activity); PRP (Properties); PUR (Purification or recovery); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(chimeric immunomodulatory compds. comprising nucleic acids linked through dendrimer or polysaccharide spacer and antigen for treating allergy, infection or cancer)

L2 ANSWER 7 OF 7 HCAPLUS COPYRIGHT 2004 ACS on STN

ED Entered STN: 05 Jul 2002

ACCESSION NUMBER: 2002:504922 HCAPLUS

DOCUMENT NUMBER: 137:73254

TITLE: Immunomodulatory oligonucleotides containing immunostimulatory sequences for treatment of disorders associated with a Th2-type immune response

INVENTOR(S): Fearon, Karen L.; Dina, Dino

PATENT ASSIGNEE(S): Dynavax Technologies Corporation, USA

SOURCE: PCT Int. Appl., 95 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002052002	A2	20020704	WO 2001-US50821	20011227
WO 2002052002	A3	20030904		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE,

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CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT,  
SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,  
SN, TD, TG

US 2003049266 A1 20030313 US 2001-33243 20011227

EP 1364010 A2 20031126 EP 2001-991610 20011227

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,  
PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

PRIORITY APPLN. INFO.: US 2000-258675P P 20001227  
WO 2001-US50821 W 20011227

OTHER SOURCE(S): MARPAT 137:73254

AB The invention provides immunomodulatory polynucleotides and methods for immunomodulation of individuals using the immunomodulatory polynucleotides. Each immunomodulatory polynucleotide comprises at least one immunostimulatory sequence (ISS). Thus, administration of 5'-tgactgtgaazgttzgagatga-3' (where z = 5-bromocytosine) in conjunction with hepatitis B surface antigen (HBsAg) to baboons resulted in increased titers of anti-HBsAg antibodies as compared to administration of HBsAg alone or to administration of HBsAg with a non-ISS oligonucleotide. Complexation of immunomodulatory polynucleotides with cationic poly(lactic acid, glycolic acid) microspheres in a human PBMC assay showed significantly enhanced induction of interferon  $\alpha$  and interferon  $\gamma$  in comparison to the polynucleotides alone.

IT 439891-56-2

RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
(immunomodulatory oligonucleotides containing immunostimulatory sequences for treatment of disorders associated with a Th2-type immune response)

E1 THROUGH E25 ASSIGNED

FILE 'REGISTRY' ENTERED AT 08:20:19 ON 06 APR 2004

L3 25 SEA FILE=REGISTRY ABB=ON PLU=ON (439891-56-2/BI OR  
482661-47-2/BI OR 482663-36-5/BI OR 482663-37-6/BI OR  
482663-46-7/BI OR 482663-95-6/BI OR 482663-96-7/BI OR  
482663-97-8/BI OR 483382-56-5/BI OR 483382-57-6/BI OR  
483382-58-7/BI OR 483382-59-8/BI OR 483382-64-5/BI OR  
483382-65-6/BI OR 483382-66-7/BI OR 483382-67-8/BI OR  
483382-68-9/BI OR 554470-62-1/BI OR 557124-76-2/BI OR  
631925-51-4/BI OR 631925-64-9/BI OR 631926-21-1/BI OR  
631926-36-8/BI OR 640803-45-8/BI OR 662376-82-1/BI)

L4 25 L1 AND L3

L4 ANSWER 1 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 662376-82-1 REGISTRY

CN DNA, d(T-C-G-T-C-G-A-A-C-G-T-T-C-G-A-G-A-T-G-A-T) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 137: PN: WO2004014322 PAGE: 24 unclaimed sequence

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

Searcher : Shears 571-272-2528

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HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 140:198064

L4 ANSWER 2 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 640803-45-8 REGISTRY  
CN DNA, d(P-thio)(T-C-G-T-C-G-A-A-C-G-T-T-C-G-A-G-A-T-G-A-T) (9CI) (CA INDEX NAME)  
CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 140:75912

L4 ANSWER 3 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 631926-36-8 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 3'-[O-(6-aminohexyl) hydrogen phosphorothioate] (9CI) (CA INDEX NAME)  
CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 140:26897

L4 ANSWER 4 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 631926-21-1 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 3'-[O-(14-hydroxy-5-mercapto-5-oxido-4,6-dioxo-10,11-dithia-5-phosphatetradec-1-yl) hydrogen phosphorothioate] (9CI) (CA INDEX NAME)  
CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t

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HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 140:26897

L4 ANSWER 5 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 631925-64-9 REGISTRY

CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,3-propanediyl oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,3-propanediyl oxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 5'-[O-(6-mercaptohexyl) hydrogen phosphorothioate] (9CI) (CA INDEX NAME)

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t

=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 140:26897

L4 ANSWER 6 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 631925-51-4 REGISTRY

CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,2-ethanediyl oxy-1,2-ethanediyl oxy-1,2-ethanediyl oxy-1,2-ethanediyl oxy-1,2-ethanediyl oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediyl oxy-1,2-ethanediyl oxy-1,2-ethanediyl oxy-1,2-ethanediyl oxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T) (9CI) (CA INDEX NAME)

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t

=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 140:26897

L4 ANSWER 7 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 557124-76-2 REGISTRY

CN DNA, d(P-thio) (T-C-G-T-C-G-A-A-C-G-T-T-C-G-A-G-A-T-G-A-T) (9CI) (CA INDEX NAME)

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t

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HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 139:99590

L4 ANSWER 8 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 554470-62-1 REGISTRY  
CN DNA, d(P-thio)(T-C-G-T-C-G-A-A-C-G-T-T-C-G-A-G-A-T-G-A-T) (9CI) (CA  
INDEX NAME)  
CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 139:83598

L4 ANSWER 9 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 483382-68-9 REGISTRY  
CN Poly(oxy-1,2-ethanediyl),  $\alpha$ -hydro- $\omega$ -hydroxy-, ester with  
2'-deoxy-P-thioadenylyl-(3' $\rightarrow$ 5')-2'-deoxy-P-thiocytidylyl-  
(3' $\rightarrow$ 5')-2'-deoxy-P-thioguanilyl-(3' $\rightarrow$ 5')-P-  
thiothymidylyl-(3' $\rightarrow$ 5')-P-thiothymidylyl-(3' $\rightarrow$ 5')-2'-  
deoxy-P-thiocytidylyl-(3' $\rightarrow$ 5')-2'-deoxyguanosine  
3',5'-bis(dihydrogen phosphorothioate), 2'-deoxy-P-thioadenylyl-  
(3' $\rightarrow$ 5')-2'-deoxy-P-thioguanilyl-(3' $\rightarrow$ 5')-2'-deoxy-P-  
thioadenylyl-(3' $\rightarrow$ 5')-P-thiothymidylyl-(3' $\rightarrow$ 5')-2'-deoxy-  
P-thioguanilyl-(3' $\rightarrow$ 5')-2'-deoxy-P-thioadenylyl-(3' $\rightarrow$ 5')-  
thymidine 5'-(dihydrogen phosphorothioate) and P-thiothymidylyl-  
(3' $\rightarrow$ 5')-2'-deoxy-P-thiocytidylyl-(3' $\rightarrow$ 5')-2'-deoxy-P-  
thioguanilyl-(3' $\rightarrow$ 5')-P-thiothymidylyl-(3' $\rightarrow$ 5')-2'-deoxy-  
P-thiocytidylyl-(3' $\rightarrow$ 5')-2'-deoxy-P-thioguanilyl-  
(3' $\rightarrow$ 5')-2'-deoxyadenosine 3'-(dihydrogen phosphorothioate)  
(2:1:1:1) (9CI) (CA INDEX NAME)  
CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 10 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 483382-67-8 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-G-

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10/033243

sp-A-sp-T-sp-G-sp-A-sp-T) (9CI) (CA INDEX NAME)  
CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 11 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 483382-66-7 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,6-hexanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,6-hexanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T) (9CI) (CA INDEX NAME)

CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 12 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 483382-65-6 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,4-butanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,4-butanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T) (9CI) (CA INDEX NAME)

CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 13 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 483382-64-5 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,3-propanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,3-propanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 5'-[O-[6-[(6-hydroxyhexyl)dithio]hexyl]hydrogen phosphorothioate] (9CI) (CA INDEX NAME)

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CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 14 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 483382-59-8 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 3'-[O-(3-mercaptopropyl) hydrogen phosphorothioate] (9CI) (CA INDEX NAME)

CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 15 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 483382-58-7 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 3'-[O-[3-(3-mercaptopropoxy)propyl] hydrogen phosphorothioate] (9CI) (CA INDEX NAME)

CI MAN  
SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 16 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
RN 483382-57-6 REGISTRY  
CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-

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ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinyl  
 idene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-  
 G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-  
 ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-  
 1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-  
 sp-A-sp-T), 5'-[O-(6-mercaptohexyl) hydrogen phosphorothioate] (9CI)  
 (CA INDEX NAME)

CI MAN  
 SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
 ===== =

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 17 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
 RN 483382-56-5 REGISTRY  
 CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)o  
 xy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-  
 ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinyl  
 idene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-  
 G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-  
 ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-  
 1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-  
 sp-A-sp-T), 5'-[O-[6-[(6-hydroxyhexyl)dithio]hexyl] hydrogen  
 phosphorothioate] (9CI) (CA INDEX NAME)

CI MAN  
 SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
 ===== =

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 18 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN  
 RN 482663-97-8 REGISTRY  
 CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)o  
 xy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-  
 ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinyl  
 idene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-  
 sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,2-  
 ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-  
 1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-  
 G-sp-A-sp-T-sp-G-sp-A-sp-T) (9CI) (CA INDEX NAME)

CI MAN  
 SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
 ===== =

HITS AT: 1-21

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## \*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 19 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 482663-96-7 REGISTRY

CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy-1,2-ethanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 3'-[O-[3-[(3-hydroxypropyl)dithio]propyl] hydrogen phosphorothioate] (9CI) (CA INDEX NAME)

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t

=====

HITS AT: 1-21

## \*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 20 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 482663-95-6 REGISTRY

CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 3'-[O-[3-[(3-hydroxypropyl)dithio]propoxy]propyl] hydrogen phosphorothioate] (9CI) (CA INDEX NAME)

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t

=====

HITS AT: 1-21

## \*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 21 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 482663-46-7 REGISTRY

CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,3-propanediylloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 5'-[O-(6-mercaptohexyl) hydrogen phosphorothioate] (9CI) (CA INDEX NAME)

CI MAN

SQL 21

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SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 22 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 482663-37-6 REGISTRY

CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,3-propanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,3-propanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T) (9CI) (CA INDEX NAME)

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 23 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 482663-36-5 REGISTRY

CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T) (9CI) (CA INDEX NAME)

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t  
=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 138:88635

L4 ANSWER 24 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 482661-47-2 REGISTRY

CN DNA, d(T-sp-C-sp-G-sp-T-sp-C-sp-G-sp-A[oxy(mercaptophosphinylidene)oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-C-sp-G-sp-T-sp-T-sp-C-sp-G[oxy(mercaptophosphinylidene)oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy-1,2-ethanediyl]oxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T) (9CI) (CA INDEX NAME)

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1,2-ethanediyloxy(mercaptophosphinylidene)oxy]A-sp-G-sp-A-sp-T-sp-G-sp-A-sp-T), 3',3''',3''''-[O,O',O''-[nitrilotris[2,1-ethanediyl(2,5-dioxo-1,3-pyrrolidinediyl)thio-3,1-propanediyl]] tris(hydrogen phosphorothioate)] (9CI) (CA INDEX NAME)

CI MAN

SQL 63,21,21,21

SEQ 1 tcgtcgaacg ttcgagatga t

=====

HITS AT: 1-21

SEQ 1 tcgtcgaacg ttcgagatga t

=====

HITS AT: 1-21

SEQ 1 tcgtcgaacg ttcgagatga t

=====

HITS AT: 1-21

REFERENCE 1: 138:88635

L4 ANSWER 25 OF 25 REGISTRY COPYRIGHT 2004 ACS on STN

RN 439891-56-2 REGISTRY

CN DNA, d(T-C-G-T-C-G-A-A-C-G-T-T-C-G-A-G-A-T-G-A-T) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 132: PN: WO02052002 SEQID: 132 claimed DNA

CI MAN

SQL 21

SEQ 1 tcgtcgaacg ttcgagatga t

=====

HITS AT: 1-21

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

REFERENCE 1: 137:73254

FILE 'HOME' ENTERED AT 08:20:41 ON 06 APR 2004

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Gencore version 5.1.6  
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## OM nucleic - nucleic search, using sw model

Run on: April 5, 2004, 21:36:48 ; Search time 2880 Seconds

(without alignments)  
316.043 Million cell updates/sec

Title: US-10-033-243-132

Sequence: 1 tcgtcgaaagtcgagatgat 21

Scoring table: IDENTITY NUC  
Gapop 10.0, Gapext 1.0

Searched: 3470272 seqs, 2167151695 residues

Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database:

```
GenEmbl.*
1: gb_ba:*
2: gb_hcg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vl:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_ov:*
22: em_pat:*
23: em_ph:*
24: em_pl:*
25: em_ro:*
26: em_sts:*
27: em_un:*
28: em_vl:*
29: em_hcg_hum:*
30: em_hcg_inv:*
31: em_hcg_other:*
32: em_hcg_mus:*
33: em_hcg_pln:*
34: em_hcg_rod:*
35: em_hcg_vrt:*
36: em_hcg_vrt:*
37: em_hcg_vrt:*
38: em_hcg_vrt:*
39: em_hcg_hum:*
40: em_hcg_mus:*
41: em_hcg_other:*
```

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	21	100.0	21	6	AX592442
2	19	90.5	19	6	AX592329
3	19	90.5	22	6	AX592340
4	18	85.7	19	6	AX592334
5	17.8	84.8	164921	8	AF022186
6	17.4	82.9	19	6	AX592366
7	17.4	82.9	19	6	AX592367
8	17	81.0	19	6	AX592333
9	16.8	80.0	658	3	TRE512595
10	16.8	80.0	3041	3	AF257638
11	16.8	80.0	3047	3	AF257637
12	16.8	80.0	3047	3	AF257639
13	16.8	80.0	3047	3	AF257640
14	16.8	80.0	3047	3	AF257641
15	16.8	80.0	3047	3	AF257643
16	16.8	80.0	3047	3	AF257646
17	16.8	80.0	3050	3	AF257642
18	16.8	80.0	3050	3	AF257644
19	16.8	80.0	3050	3	AF257645
20	16.8	80.0	3050	3	AF257647
21	16.8	80.0	3050	3	AF257648
22	16.8	80.0	3050	3	AF257649
23	16.8	80.0	146568	2	AC141727
24	16.4	78.1	18	6	AX592324
25	16.4	78.1	23	6	BD233619
26	16.4	78.1	23	6	AR352575
27	16.4	78.1	23	6	AX083677
28	16.4	78.1	23	6	AX148638
29	16.4	78.1	23	6	AX250703
30	16.4	78.1	23	6	AX252293
31	16.4	78.1	23	6	AX252511
32	16.4	78.1	23	6	AX252522
33	16.4	78.1	23	6	AX252936
34	16.4	78.1	23	6	AX253115
35	16.4	78.1	23	6	AX253125
36	16.4	78.1	416	8	ATH528591
37	16.4	78.1	68415	8	AC005310
38	16.4	78.1	73921	8	AB024024
39	16.2	77.1	1687	8	AK100785
40	16.2	77.1	2064	5	AF069993
41	16.2	77.1	3135	6	AR346964
42	16.2	77.1	3960	8	AF152552
43	16.2	77.1	3989	4	BTU61233
44	16.2	77.1	7749	2	AC014752
45	16.2	77.1	19226	14	CTU56902

## ALIGNMENTS

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RESULT 1
LOCUS AX592442
DEFINITION Sequence 132 from Patent WO02032002.
ACCESSION AX592442
VERSION AX592442.1 GI:27950544
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1
AUTHORS Fearon,K.L. and Dina,D.
TITLE Immunomodulatory polynucleotides and methods of using the same
JOURNML Patent: WO 02052002-A 132 04-JUL-2002;
DynaVax Technologies Corporation (US)
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## FEATURES

Location/Qualifiers  
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/mol\_type="unassigned DNA"  
/db\_xref="taxon:32630"  
/note="Polynucleotide containing CG"

## ORIGIN

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Best Local Similarity 100.0%; Pred. No. 0.6;  
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTGAACGTTCCGAGATG 21  
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1 TCGTGAACGTTCCGAGATG 21

## RESULT 2

AX592329 19 bp DNA linear PAT 27-JAN-2003  
LOCUS AX592329  
DEFINITION Sequence 19 from Patent WO02052002.  
ACCESSION AX592329  
VERSION AX592329.1 GI:27950431  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS Fearon,K.L. and Dina,D.  
TITLE Immunomodulatory polynucleotides and methods of using the same  
JOURNAL Patent: WO 02052002-A 19 04-JUL-2002;  
Dynavax Technologies Corporation (US)  
Location/Qualifiers

## FEATURES

1..19  
/organism="synthetic construct"  
/mol\_type="unassigned DNA"  
/db\_xref="taxon:32630"  
/note="Polynucleotide containing CG"

## ORIGIN

Query Match 90.5%; Score 19; DB 6; Length 19;  
Best Local Similarity 100.0%; Pred. No. 8.6;  
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTGAACGTTCCGAGATG 19  
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## RESULT 3

AX592340 22 bp DNA linear PAT 27-JAN-2003  
LOCUS AX592340  
DEFINITION Sequence 30 from Patent WO02052002.  
ACCESSION AX592340  
VERSION AX592340.1 GI:27950442  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS Fearon,K.L. and Dina,D.  
TITLE Immunomodulatory polynucleotides and methods of using the same  
JOURNAL Patent: WO 02052002-A 30 04-JUL-2002;  
Dynavax Technologies Corporation (US)  
Location/Qualifiers

## FEATURES

1..22  
/organism="synthetic construct"  
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/note="Polynucleotide containing CG"

## ORIGIN

Query Match 90.5%; Score 19; DB 6; Length 22;

Best Local Similarity 100.0%; Pred. No. 8.7;  
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTGAACGTTCCGAGATG 19  
|||||  
4 TCGTGAACGTTCCGAGATG 22

## RESULT 4

AX592334 19 bp DNA linear PAT 27-JAN-2003  
LOCUS AX592334  
DEFINITION Sequence 24 from Patent WO02052002.  
ACCESSION AX592334  
VERSION AX592334.1 GI:27950436  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS Fearon,K.L. and Dina,D.  
TITLE Immunomodulatory polynucleotides and methods of using the same  
JOURNAL Patent: WO 02052002-A 24 04-JUL-2002;  
Dynavax Technologies Corporation (US)  
Location/Qualifiers

## FEATURES

1..19  
/organism="synthetic construct"  
/mol\_type="unassigned DNA"  
/db\_xref="taxon:32630"  
/note="Polynucleotide containing CG"  
misc\_feature 5  
/note="n = 5-bromocytosine"

## ORIGIN

Query Match 85.7%; Score 18; DB 6; Length 19;  
Best Local Similarity 94.7%; Pred. No. 33;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTGAACGTTCCGAGATG 19  
|||||  
1 TCGTGAACGTTCCGAGATG 19

## RESULT 5

AF022186 164921 bp DNA circular PLN 14-DEC-2000  
LOCUS AF022186  
DEFINITION Cyanidium caldarium strain RK1 chloroplast, complete genome.  
ACCESSION AF022186 Z36235 Z70297  
VERSION AF022186.2 GI:6466296  
KEYWORDS  
SOURCE chloroplast Cyanidium caldarium  
ORGANISM Cyanidium caldarium

Eukaryota; Rhodophyta; Bangiophyceae; Porphyridiales;  
Porphyridiaceae; Cyanidium.

## REFERENCE

1 (bases 130696 to 133664)  
Vogel,H., Fischer,S. and Valentín,K.  
A model for the evolution of the plastid sec apparatus inferred  
from secY gene phylogeny

## JOURNAL

Plant Mol. Biol. 32 (4), 685-692 (1996)  
MEDLINE 97134960  
PUBMED 8980520  
2 (bases 1 to 164921)  
Glockner,G., Rosenthal A. and Valentín,K.  
The structure and gene repertoire of an ancient red algal plastid  
genome

## JOURNAL

J. Mol. Evol. 51 (4), 382-390 (2000)  
MEDLINE 20496959  
PUBMED 11040290  
3 (bases 46857 to 47851)  
Valentin,K.  
Direct Submission

## JOURNAL

Submitted (22-MAR-1996) Institute for Plant Physiology, Justus  
Liebig University, Heinrich Buff Ring 58-62, Gießen 35392, Germany  
REFERENCE 4 (bases 28701 to 75580)



AUTHORS Gloeckner,G., Rosenthal,A. and Valentini,K.  
TITLE Direct Submission  
JOURNAL Submitted (02-SEP-1997) Department of Genome Analysis, IMB Jena,  
Beutenbergstr.11, Jena 07745, Germany  
REFERENCE 5 (bases 1 to 164921)  
AUTHORS Gloeckner,G., Rosenthal,A. and Valentini,K.  
TITLE Direct Submission  
JOURNAL Submitted (18-NOV-1999) Genome Analysis, Institute for Molecular  
Biotechnology, Beutenbergstrasse 11, Jena 07745, Germany  
6 (bases 13066 to 132364)  
AUTHORS Vogel,H., Fischer,S. and Valentini,K.  
TITLE Direct Submission  
JOURNAL Submitted (18-NOV-1999) Institute for Plant Physiology, Justus  
Liebig University, Heinrich Buff Ring 58-62, Giessen 35392, Germany  
On or before Nov 23, 1999 this sequence version replaced gi:529651,  
gi:1240002, gi:2465730.  
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339..477  
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580..780  
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KEYWORDS	synthetic construct
SOURCE	synthetic construct
ORGANISM	artificial sequences.
REFERENCE	1 Pearson, K.L. and Dina, D. Immunomodulatory polynucleotides and methods of using the same
AUTHORS	Patent: WO 02052002-A 57 04-Jul-2002;
TITLE	Dynavax Technologies Corporation (US)
JOURNAL	Location/Qualifiers
FEATURES	1..19
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Best Local Similarity	94.7%; Pred. No. 73;
Matches	18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY	1 TCGTGAACGTTGCAGATG 19
Db	1 TCGTGAACGTTGCAGATG 19
RESULT 8	
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LOCUS	AX592333
DEFINITION	Sequence 23 from Patent WO02052002.
ACCESSION	AX592333
VERSION	AX592333.1 GI:27950435
KEYWORDS	
SOURCE	synthetic construct
ORGANISM	synthetic construct
REFERENCE	1 artificial sequences.
AUTHORS	1 Pearson, K.L. and Dina, D. Immunomodulatory polynucleotides and methods of using the same
TITLE	Patent: WO 02052002-A 23 04-Jul-2002;
JOURNAL	Dynavax Technologies Corporation (US)
FEATURES	Location/Qualifiers
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	5
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ORIGIN	
Query Match	81.0%; Score 17; DB 6; Length 19;
Best Local Similarity	89.5%; Pred. No. 1.2e+02;
Matches	17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
OY	1 TCGTGAACGTTGCAGATG 19
Db	1 TNGTGAACGTTGCAGATG 19
RESULT 9	
TR512595	658 bp DNA linear INV 18-AUG-2003
LOCUS	TR512595
DEFINITION	Timarcha recticollis 5.8S rRNA gene (partial), 26S rRNA (partial)
ACCESSION	U512595
VERSION	U512595.1 GI:3394511
KEYWORDS	26S ribosomal RNA; 5.8S ribosomal RNA; 5.8S rRNA
SOURCE	gene, internal transcribed spacer 2; ITS2.
ORGANISM	Timarcha recticollis
	Timarcha recticollis

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HEFGDTGQDLDFIRNEFDRLLBTD  
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VTNRAAMLVASGVSC.LIDMR

EEFVITQDSCGVGAIMAGMAH

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VS E D S G R G A A L V A A T A V Q A K

gth 3041;

models 0; Gaps 0;

near INV 09-OCT-2000

77-368170YAH DUM T 7-

Acta; Pterygota;  
Muscomorpha;

our hexokinases in

State University of

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SKL"

ORIGIN
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Best Local Similarity 90.0%; Pred. No. 3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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|||||
156 CGTCGACGATCGATGAT 137

Db

RESULT 12
AF257639/c      3047 bp      DNA      linear      INV 09-OCT-2000
LOCUS
DEFINITION
Drosophila simulans strain CT96_5s hexokinase-t1 and hexokinase-t2
genes, complete cds.
ACCESSION
AF257639
VERSION
AF257639.1 GI:10765246
KEYWORDS
SOURCE
Drosophila simulans
ORGANISM
Drosophila simulans
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 3047)
Duvernell, D.D. and Eanes, W.F.
Contrasting molecular population genetics of four hexokinases in
Drosophila melanogaster and Drosophila simulans
Genetics (2000) In press
2 (bases 1 to 3047)
Duvernell, D.D. and Eanes, W.F.
Direct Submission
Submitted (19-APR-2000) Ecology and Evolution, State University of
New York, Stony Brook, NY 11794, USA
JOURNAL
TITLES
Location/Qualifiers
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FEATURES
source
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SKL"

ORIGIN
Query Match      80.0%; Score 16.8; DB 3; Length 3047;
Best Local Similarity 90.0%; Pred. No. 3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY
2 CGTCGACGTCGAGATGAT 21
|||||
156 CGTCGACGATCGATGAT 137

Db

RESULT 13
AF257640/c      3047 bp      DNA      linear      INV 09-OCT-2000
LOCUS
DEFINITION
Drosophila simulans strain CT96_6s hexokinase-t1 and hexokinase-t2
genes, complete cds.
ACCESSION
AF257640
VERSION
AF257640.1 GI:10765249
KEYWORDS
SOURCE
Drosophila simulans
ORGANISM
Drosophila simulans
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 3047)
Duvernell, D.D. and Eanes, W.F.
Contrasting molecular population genetics of four hexokinases in
Drosophila melanogaster and Drosophila simulans
Genetics (2000) In press
2 (bases 1 to 3047)
Duvernell, D.D. and Eanes, W.F.
Direct Submission
Submitted (19-APR-2000) Ecology and Evolution, State University of
New York, Stony Brook, NY 11794, USA
JOURNAL
TITLES
Location/Qualifiers
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ACCESSION	AF257643				

KEYWORDS	SOURCE	ORGANISM
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	Drosophila simulans	
Eukaryota: Metazoa: Arthropoda: Insecta: Pterygota		

1, D.D. and Eanes, W.F.  
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**TITLE** Contrasting molecular population genetics of four hexokinases in *Drosophila melanogaster* and *Drosophila simulans*

**JOURNAL REFERENCE** Genetics (2000) In Press

**AUTHORS** 2 (baaes 1 to 3047)

**TITLE** Duvernell, D.D. and Eanes, W.F.

**JOURNAL** Direct Submission

**FEATURES** Submitted (19-Apr-2000) Ecology and Evolution, State University of New York, Stony Brook, NY 11794, USA

**LOCATION/QUALIFIERS** Location/Qualifiers

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SOURCE Drosophila simulans  
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REFERENCE  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE 1 (bases 1 to 3047)  
JOURNAL Contrasting molecular population genetics of four hexokinases in  
Drosophila melanogaster and Drosophila simulans  
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TITLE Direct Submission  
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Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
Ephydriidae; Drosophilidae; Drosophila.  
REFERENCE  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE 1 (bases 1 to 3050)  
JOURNAL Contrasting molecular population genetics of four hexokinases in  
Drosophila melanogaster and Drosophila simulans  
GENETICS (2000) In press  
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REFERENCE  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE Direct Submission  
JOURNAL Submitted (19-APR-2000) Ecology and Evolution, State University of  
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## ORIGIN

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Db 156 CGTCGACGATCGTATGAT 137

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VERSION AF257644.1  
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SOURCE Drosophila simulans  
ORGANISM Drosophila simulans  
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Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
Ephydroidae; Drosophilidae; Drosophila.  
REFERENCE  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE Contrasting molecular population genetics of four hexokinases in  
Drosophila melanogaster and Drosophila simulans  
JOURNAL Genetics (2000) In press  
REFERENCE  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE Direct Submission  
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Db 156 CGTCGACGATCGTATGAT 137

RESULT 19  
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LOCUS Drosophila simulans strain GA96\_4a hexokinase-t1 and hexokinase-t2  
DEFINITION genes, complete cds.  
ACCESSION AF257645  
VERSION AF257645.1 GI:10765264  
KEYWORDS  
SOURCE Drosophila simulans  
ORGANISM Drosophila simulans  
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
Ephydroidae; Drosophilidae; Drosophila.  
REFERENCE  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE Contrasting molecular population genetics of four hexokinases in  
Drosophila melanogaster and Drosophila simulans  
JOURNAL Genetics (2000) In press  
REFERENCE  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
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genes, complete cds.  
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VERSION AF257647.1 GI:10765270  
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SOURCE Drosophila simulans  
ORGANISM Drosophila simulans  
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Ephydroidea; Drosophilidae; Drosophila.  
REFERENCE 1 (bases 1 to 3050)  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE Contrasting molecular population genetics of four hexokinases in  
Drosophila melanogaster and Drosophila simulans  
JOURNAL Genetics (2000) In press  
2 (bases 1 to 3050)  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
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Best Local Similarity 90.0%; Pred. No. 3e+02;  
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REFERENCE 1 (bases 1 to 3050)  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE Contrasting molecular population genetics of four hexokinases in  
Drosophila melanogaster and Drosophila simulans  
JOURNAL Genetics (2000) In press  
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AUTHORS Duvernell,D.D. and Eanes,W.F.  
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SKL"

ORIGIN

Query Match 80.0%; Score 16.8; DB 3; Length 3050;  
Beet Local Similarity 90.0%; Pred. No. 3e+02;  
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CGTGAAGCTTCGAGATGAT 21  
Db 156 CGTGAAGCTTCGATGAT 137

RESULT 22  
LOCUS AF257649 3050 bp DNA linear INV 09-OCT-2000  
DEFINITION Drosophila simulans strain HFL97\_48 hexokinase-t2  
genes, complete cds.  
ACCESSION AF257649  
VERSION AF257649.1 GI:10765276  
KEYWORDS  
SOURCE Drosophila simulans  
ORGANISM Drosophila simulans  
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
Ephydroidea; Drosophilidae; Drosophila.  
REFERENCE 1 (bases 1 to 3050)  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE Contrasting molecular population genetics of four hexokinases in  
Drosophila melanogaster and Drosophila simulans  
JOURNAL Genetics (2000) In press  
RECORD 2 (bases 1 to 3050)  
AUTHORS Duvernell,D.D. and Eanes,W.F.  
TITLE Direct Submission  
JOURNAL Submitted (19-Apr-2000) Ecology and Evolution, State University of  
New York, Stony Brook, NY 11794, USA  
FEATURES  
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/db\_xref="taxon:7240"  
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/db\_xref="GI:10765277"  
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WTERGAGAGIGKDVQVLLRDALAKPEISVDVMTINVAGSGLATLCAWAPDRIGL  
IMSTIANSCYVERECETTBGDEFRRMLMTINSMAHFGDQGLDFINERDRLLD  
SINGSTRYERKESVFCALCMGELVRIILVILMKAGAFADRDYIGIOWKLDVSLIEI  
VSDPPGVYTKAOEWMDEKFRIRHCKERDLAALKYICDVTNRAAMLVAGVSCLDLDRM  
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LMSGKGLPDLFLAESLSEFCHTGLENESLPLGTFSPFLQOGLSGIIVAMKGF  
SCEGVGKNVVLLOEALDRRGDIKINTVALINTVGTLMGCAFPMPCRIGLIVGTG  
SNACVTEKTVNAECFEGYQTSRPSMTINCENMGAFDNGVUEPRTSDKIVDKYTPN  
PGKOTFEKICISGMWGLVRLVLDIMAKFMPHGIISEKIOERMSFTAYISDVESD  
APGERNCNKVLSLGIIGCEPDKELRYICEAVSSRSKALCAGGLVTTINKMINE  
VVIIGIDSVYRFHFKYHDMLOHMKKLLKPKVKEELIVSEDSGSGAALVAATAVOAK  
SKL"

ORIGIN

Query Match 80.0%; Score 16.8; DB 3; Length 3050;  
Beet Local Similarity 90.0%; Pred. No. 3e+02;  
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CGTGAAGCTTCGAGATGAT 21  
Db 156 CGTGAAGCTTCGATGAT 137

RESULT 23  
LOCUS AC141727/c 146568 bp DNA linear HTG 19-MAR-2003  
DEFINITION Apis mellifera clone CH224-5763, WORKING DRAFT SEQUENCE, 25  
unordered pieces.  
ACCESSION AC141727  
VERSION AC141727.1 GI:29123911  
KEYWORDS HTGS PHASE1, HTGS DRAFT.  
SOURCE Apis mellifera (honeybee)  
ORGANISM Apis mellifera  
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
Neoptera; Endopterygota; Hymenoptera; Apocrita; Aculeata; Apoidea;  
Apidae; Apis.  
REFERENCE 1 (bases 1 to 146568)  
AUTHORS Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-ozman,F.R., Allen,C.,  
Alldbrooks,S.L., Amaralunga,H.C., Are,J.R., Ayala,M., Banks,T.,  
Barbarta,J., Benton,J., Blum,K., Blankenburg,K., Bonnin,D.,  
Bouck,J., Bowle,S., Brileva,M., Brown,E., Brown,M., Bryant,N.P.,  
Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C.,  
Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D.,  
Chen,G., Chen,R., Chen,Z., Chowdhury,I., Christopoulos,C.,  
Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R.,  
Dayila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A.,  
Delaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinn,H.H.,  
Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Durbin,K.J.,  
Earnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Escoto,M.,  
Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P.,  
Gabisi,J., Gao,U., Garcia,A., Garner,T., Garza,N., Gill,R.,  
Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K.,  
Harris,C., Harris,K., Hart,M., Havlak,P., Hawes,A., Hernandez,J.,  
Hernandez,O., Hodgson,A., Hognes,M., Holloway,C., Hollins,B.,  
Honsi,F., Howard,S., Huber,J., Hulyk,S., Hume,J., Jackson,L.E.,  
Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudah,S.,  
Karleson,E., Kelly,S., Khan,U., King,L., Korvah,U., Kovar,C.,  
Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C., Lewis,L.,  
Li,J., Li,Z., Lichtarge,O., Lieu,A., Liu,B., Liu,W., Louised,H.,  
Lohado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J.,  
Maheshwari,M., Mapua,P., Martin,R., Martinale,A., Martinez,E.,  
Massey,E., Mawhinney,E., McLeod,M.P., Meddor,M., Mei,G., Metker,M.,  
Miner,G., Miner,Z., Mitchell,T., Mohabbat,K., Morgan,M., Morris,S.,

Moser, M., Neal, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokoko, S., Ogun, M., Okunolu, G., Oregan, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L., Quiles, M., Ren, Y., Rivers, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savery, G., Scherer, S., Scott, G., Shen, H., Shoshkari, N., Sison, I., Sodergren, E., Sonaike, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Taber, P., Tamerisa, A., Tamerisa, K., Tang, H., Taney, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S., Usmani, K., Vaequez, L., Vera, V., Villalon, D., Vinson, R., Wang, S., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S., Williams, G., Williamson, A., Wleczek, R., Wooden, S., Worley, K., Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Nelson, D., Weinstein, G. and Gibbs, R.

Unpublished  
2 (bases 1 to 146568)  
Worley, K.C.

Direct Submission  
Submitted (19-MAR-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

----- Genome Center -----  
Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: hgsc-help@bcm.tmc.edu

----- Project Information -----  
Center project name: AMBZ  
Center clone name: CH224-57G3  
----- Summary Statistics -----  
Sequencing vector: Plasmid;  
Chemistry: Dye-terminator Big Dye 100% of reads  
Assembly program: Phrap; version 0.990329  
Consensus quality: 132495 bases at least Q40  
Consensus quality: 137202 bases at least Q30  
Consensus quality: 140042 bases at least Q20  
Estimated insert size: 137666; sum-of-contigs estimation  
Quality coverage: 4x in Q20 bases; sum-of-contigs estimation

----- NOTE: Estimated insert size may differ from sequence length -----  
(see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 25 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1 1456: contig of 1456 bp in length  
\* 1457 1556: gap of unknown length  
\* 1557 2731: contig of 1175 bp in length  
\* 2732 2831: gap of unknown length  
\* 2832 4180: contig of 1349 bp in length  
\* 4181 4280: gap of unknown length  
\* 4281 5835: contig of 1555 bp in length  
\* 5836 5935: gap of unknown length  
\* 5936 7358: contig of 1423 bp in length  
\* 7359 7458: gap of unknown length  
\* 7459 8816: contig of 1358 bp in length  
\* 8817 8916: gap of unknown length  
\* 8917 10936: contig of 2020 bp in length  
\* 10937 11036: gap of unknown length  
\* 11037 12381: contig of 1345 bp in length  
\* 12382 12481: gap of unknown length  
\* 12482 15062: contig of 2581 bp in length  
\* 15063 15162: gap of unknown length  
\* 15163 18933: contig of 3771 bp in length  
\* 18934 19033: gap of unknown length  
\* 19034 23106: contig of 4073 bp in length  
\* 23107 23206: gap of unknown length  
\* 23207 27707: contig of 4501 bp in length

27708 27807: gap of unknown length  
\* 27808 33059: contig of 5252 bp in length  
\* 33060 33159: gap of unknown length  
\* 33160 38669: contig of 5510 bp in length  
\* 38670 45437: gap of unknown length  
\* 45437 45536: gap of unknown length  
\* 45537 52247: contig of 6711 bp in length  
\* 52248 52347: gap of unknown length  
\* 52348 52944: contig of 6897 bp in length  
\* 52945 59344: gap of unknown length  
\* 59345 66497: contig of 7153 bp in length  
\* 66498 66597: gap of unknown length  
\* 66598 74127: contig of 7530 bp in length  
\* 74128 74227: gap of unknown length  
\* 74228 81822: contig of 7495 bp in length  
\* 81823 81822: gap of unknown length  
\* 81823 92875: contig of 11053 bp in length  
\* 92876 92975: gap of unknown length  
\* 92976 101243: contig of 8268 bp in length  
\* 101244 101343: gap of unknown length  
\* 101344 113869: contig of 12526 bp in length  
\* 113870 113969: gap of unknown length  
\* 113970 128447: contig of 14478 bp in length  
\* 128448 128547: gap of unknown length  
\* 128548 146568: contig of 18021 bp in length.

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ORIGIN  
Query Match 80.0%; Score 16.8; DB 2; Length 146568;  
Best Local Similarity 90.0%; Pred. No. 4.7e+02;  
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 CGTCGACGTCGAGATGAT 21  
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Db 135270 CGTCGACGTCGAGACGTT 135251

RESULT 24  
AX592324 18 bp DNA linear PAT 27-JAN-2003  
DEFINITION Sequence 14 from Patent WO02052002.  
ACCESSION AX592324  
VERSION AX592324.1 GI:27950426  
KEYWORDS  
SOURCE  
ORGANISM  
synthetic construct  
artificial sequences.  
REFERENCE  
1  
Featon, K.L. and Dina, D.  
Immunomodulatory polynucleotides and methods of using the same  
Patent: WO 02052002-A 14 04-JUL-2002;  
Dynavax Technologies Corporation (US)  
FEATURES  
source  
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/mol\_type="unassigned DNA"  
/db\_xref="taxon:32630"  
/note="Polynucleotide containing CG"

ORIGIN  
Query Match 78.1%; Score 16.4; DB 6; Length 18;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 CGTCGACGTCGAGATG 19  
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Db 1 CTTGCAACGTCGAGATG 18

RESULT 25  
BD233619/c  
LOCUS 23 bp DNA linear PAT 17-JUL-2003  
DEFINITION Immunostimulatory oligonucleotides, compositions thereof and methods of use thereof.  
ACCESSION BD233619  
VERSION BD233619.1 GI:33043389  
KEYWORDS JP 2002517156-A/4.  
SOURCE unidentified  
ORGANISM unidentified  
REFERENCE 1 (bases 1 to 23)  
AUTHORS Schwartz,D., Roman,M., Dina,D. and Raz,E.  
TITLE Immunostimulatory oligonucleotides, compositions thereof and methods of use thereof  
JOURNAL Patent: JP 2002517156-A 4 11-JUN-2002;  
COMMENT DYNVAX TECHNOLOGIES CORP  
OS Unidentified  
PN JP 2002517156-A/4  
PD 11-JUN-2002  
PF 05-JUN-1998 JP 1999502884  
PR 06-JUN-1997 US 60/048793  
PI DAVID SCHWARTZ, MARK ROMAN, DINO DINA, EYAL RAZ  
PC C12N15/09,A61K31/7088,A61K31/7115,A61P37/02,A61P43/00,C12Q1/68, PC C12N15/00  
CC Strandedness: Single;  
CC Topology: Linear;  
CC Immunostimulatory oligonucleotides, compositions thereof and methods of use thereof  
CC use thereof  
FH Key 1.23 Location/Qualifiers  
FT source 1.23 /organism='Unidentified'.  
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ORIGIN  
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Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGCAGATGA 20  
DB 18 GTGGAACGTTGCAGATGA 1

RESULT 26  
AR352575/c  
LOCUS 23 bp DNA linear PAT 17-AUG-2003  
DEFINITION Sequence 4 from patent US 6589940.  
ACCESSION AR352575  
VERSION AR352575.1 GI:33757826  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 23)  
AUTHORS Raz,E., Roman,M. and Dina,D.  
TITLE Immunostimulatory oligonucleotides, compositions thereof and methods of use thereof  
JOURNAL Patent: US 6589940-A 4 08-JUL-2003;  
FEATURES  
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Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGCAGATGA 20  
DB 18 GTGGAACGTTGCAGATGA 1

RESULT 27  
AX083677/c  
LOCUS 23 bp DNA linear PAT 28-FEB-2001  
DEFINITION Sequence 3 from Patent WO0112223.  
ACCESSION AX083677  
VERSION AX083677.1 GI:13185409  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS van Nest,G.  
TITLE Methods of modulating an immune response using immunostimulatory sequences and compositions for use therein  
JOURNAL Patent: WO 0112223-A 3 22-FEB-2001;  
FEATURES  
source 1.23 Location/Qualifiers  
1.23 /organism='synthetic construct'  
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/note='Synthetic construct'

ORIGIN  
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Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGCAGATGA 20  
DB 18 GTGGAACGTTGCAGATGA 1

RESULT 28  
AX148638/c  
LOCUS 23 bp DNA linear PAT 08-JUN-2001  
DEFINITION Sequence 3 from Patent WO0135991.  
ACCESSION AX148638  
VERSION AX148638.1 GI:14347256  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS Tuck,S. and van Nest,G.  
TITLE Immunomodulatory compositions containing an immunostimulatory sequence linked to antigen and methods of use thereof  
JOURNAL Patent: WO 0135991-A 3 25-MAY-2001;  
FEATURES  
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/note='Synthetic construct'

ORIGIN  
Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

RESULT 29  
AX250703/c 23 bp DNA linear PAT 05-OCT-2001  
LOCUS Sequence 3 from Patent WO0168078.  
DEFINITION AX250703  
ACCESSION AX250703  
VERSION AX250703.1 GI:15984441  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS van Nest,G.  
TITLE Methods of suppressing hepatitis virus infection using immunomodulatory polynucleotide sequences  
JOURNAL Patent: WO 0168078-A 3 20-SEP-2001;  
DynaVax Technologies Corporation (US)  
FEATURES  
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Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
Qy 3 GTCGAACGTTGAGATGA 20  
Db 18 GTCGAACGTTGAGATGA 1  
RESULT 30  
AX252293/c 23 bp DNA linear PAT 05-OCT-2001  
LOCUS AX252293  
DEFINITION Sequence 3 from Patent WO0168117.  
ACCESSION AX252293  
VERSION AX252293.1 GI:15985634  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS van Nest,G.  
TITLE Methods of reducing papillomavirus infection using immunomodulatory polynucleotide sequences  
JOURNAL Patent: WO 0168117-A 3 20-SEP-2001;  
DynaVax Technologies Corporation (US)  
FEATURES  
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Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
Qy 3 GTCGAACGTTGAGATGA 20  
Db 18 GTCGAACGTTGAGATGA 1  
RESULT 31  
AX252511/c 23 bp DNA linear PAT 05-OCT-2001  
LOCUS AX252511  
DEFINITION Sequence 3 from Patent WO0168103.  
ACCESSION AX252511  
VERSION AX252511.1 GI:15985782

KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS van Nest,G.  
TITLE Methods of ameliorating symptoms of herpes infection using immunomodulatory polynucleotide sequences  
JOURNAL Patent: WO 0168103-A 3 20-SEP-2001;  
DynaVax Technologies Corporation (US)  
FEATURES  
source 1..23  
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/note="Polynucleotide containing CG"  
ORIGIN  
Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
Qy 3 GTCGAACGTTGAGATGA 20  
Db 18 GTCGAACGTTGAGATGA 1  
RESULT 32  
AX252522/c 23 bp DNA linear PAT 05-OCT-2001  
LOCUS AX252522  
DEFINITION Sequence 3 from Patent WO0168144.  
ACCESSION AX252522  
VERSION AX252522.1 GI:15985793  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS van Nest,G. and Tuck,S.  
TITLE Biodegradable immunomodulatory formulations and methods for use thereof  
JOURNAL Patent: WO 0168144-A 3 20-SEP-2001;  
DynaVax Technologies Corporation (US)  
FEATURES  
source 1..23  
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/note="Polynucleotide containing CG"  
ORIGIN  
Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
Qy 3 GTCGAACGTTGAGATGA 20  
Db 18 GTCGAACGTTGAGATGA 1  
RESULT 33  
AX252936/c 23 bp DNA linear PAT 05-OCT-2001  
LOCUS AX252936  
DEFINITION Sequence 3 from Patent WO0168143.  
ACCESSION AX252936  
VERSION AX252936.1 GI:15986203  
KEYWORDS  
SOURCE synthetic construct  
ORGANISM synthetic construct  
REFERENCE 1  
AUTHORS van Nest,G. and Tuck,S.  
TITLE Immunomodulatory formulations and methods for use thereof

JOURNAL Patent: WO 0168143-A 3 20-SEP-2001;  
Dynavax Technologies Corporation (US)  
FEATURES  
SOURCE  
1. .23  
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/note="Polynucleotide containing CG"

ORIGIN  
Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GTCGACGCTTCGAGATGA 20  
|||  
18 GTGGAACGCTTCGAGATGA 1

Db

RESULT 34  
AX253115/c 23 bp DNA linear PAT 05-OCT-2001  
LOCUS  
DEFINITION Sequence 3 from Patent WO0168116.  
ACCESSION AX253115  
VERSION AX253115.1 GI:15986283  
KEYWORDS  
SOURCE  
ORGANISM  
synthetic construct  
synthetic construct  
artificial sequences.

REFERENCE  
1  
AUTHORS  
van Nest, G.  
TITLE  
Methods of preventing and treating respiratory viral infection using immunomodulatory polynucleotide sequences  
JOURNAL  
Dynavax Technologies Corporation (US)  
LOCATION/Qualifiers  
1. .23  
/organism="synthetic construct"  
/mol\_type="unassigned DNA"  
/db\_xref="taxon:32630"  
/note="Polynucleotide containing CG"

FEATURES  
SOURCE

ORIGIN  
Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GTCGACGCTTCGAGATGA 20  
|||  
18 GTGGAACGCTTCGAGATGA 1

Db

RESULT 35  
AX253125/c 23 bp DNA linear PAT 05-OCT-2001  
LOCUS  
DEFINITION Sequence 3 from Patent WO0168077.  
ACCESSION AX253125  
VERSION AX253125.1 GI:15986293  
KEYWORDS  
SOURCE  
ORGANISM  
synthetic construct  
synthetic construct  
artificial sequences.

REFERENCE  
1  
AUTHORS  
van Nest, G.  
TITLE  
Methods of preventing and treating viral infections using immunomodulatory polynucleotide sequences  
JOURNAL  
Dynavax Technologies Corporation (US)  
LOCATION/Qualifiers  
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FEATURES  
SOURCE

ORIGIN  
Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 2.8e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 TCGAACGCTTCGAGATGAT 21  
|||  
155 TCGAACGCTTCGAGATGAT 172

Db

RESULT 36  
ATH528591 416 bp DNA linear PLN 29-MAR-2003  
LOCUS  
DEFINITION Arabidopsis thaliana T-DNA flanking sequence, left border, clone 168C08.  
ACCESSION AJ528591  
VERSION AJ528591.1 GI:26796851  
KEYWORDS  
left border; T-DNA flanking sequence.  
SOURCE  
Arabidopsis thaliana (thale cress)  
ORGANISM  
Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE  
1  
AUTHORS  
Brunaud, V., Balzergue, S., Dubreucq, B., Aubourg, S., Samson, F., Chauvin, S., Bechtold, N., Gruaud, C., Derose, R., Pelletier, G., Lepintec, L., Caboche, M. and Lecharny, A.  
TITLE  
T-DNA integration into the Arabidopsis genome depends on sequences of pre-insertion sites  
EMBO Rep. 3 (12), 1152-1157 (2002)

JOURNAL  
MEDLINE  
22363535  
PUBMED  
12446565  
REFERENCE  
2 (bases 1 to 416)  
AUTHORS  
Balzergue, S.  
TITLE  
Direct Submission  
JOURNAL  
Submitted (21-NOV-2002) Balzergue S., UMRGV, INRA/CNRS, 2 rue Gaston Cremieux, 91057 Evry cedex, FRANCE  
COMMENT  
PCR was performed on DNA from transformants of Arabidopsis thaliana plants from INRA (Versailles). The DNA fragment(s) resulting from the PCR were directly sequenced from the left or the right border to determine the genomic sequence flanking the insertion. T-DNA derived sequences were removed. Information to order the corresponding mutant line and a link to a database providing a graphical display of the insertion site are available at <http://dbseqg.versailles.inra.fr/publiclines/>. This sequence has been generated in the framework of the French plant genomics program 'Genoplante' (<http://www.genoplante.com> and <http://genoplante-info.inbio.gen.fr>).  
Location/Qualifiers  
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1. .416  
/note="T-DNA flanking sequence  
left border"

FEATURES  
SOURCE  
misc\_feature  
1. .416  
/note="T-DNA flanking sequence  
left border"

ORIGIN  
Query Match 78.1%; Score 16.4; DB 8; Length 416;  
Best Local Similarity 94.4%; Pred. No. 4e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 TCGAACGCTTCGAGATGAT 21  
|||  
155 TCGAACGCTTCGAGATGAT 172

Db

RESULT 37

AC005310 68415 bp DNA linear PLN 27-FEB-2002  
LOCUS Arabidopsis thaliana chromosome 2 clone F19D11 map C1C06C03,  
DEFINITION complete sequence.  
ACCESSION AC005310  
VERSION AC005310.3 GI:20197315  
SOURCE HTG  
ORGANISM Arabidopsis thaliana (thale cress)  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.  
1 (bases 1 to 68415)  
Rounsley,S.D., Lin,X., Kaul,S., Shea,T.P., Fujii,C.Y., Mason,T.M.,  
Shen,M., Konning,C.M., Frazer,C.M., Somerville,C.R. and Venter,J.C.  
Unpublished  
2 (bases 1 to 68415)  
Lin,X.  
JOURNAL Direct Submission  
REFERENCE Submitted (09-MAR-2000) The Institute for Genomic Research, 9712  
AUTHORS Medical Center Dr., Rockville, MD 20850, USA  
TITLE 3 (bases 1 to 68415)  
JOURNAL Town,C.D. and Kaul,S.  
REFERENCE Direct Submission  
AUTHORS Submitted (27-FEB-2002) The Institute for Genomic Research, 9712  
JOURNAL Medical Center Dr., Rockville, MD 20850, USA, cdtowm@tigr.org  
COMMENT On Apr 18, 2002 this sequence version replaced gi:6598455.  
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KETEALIDFDIYVRANDPLTLPRLYEDFLIEIEQIALKYNALPHWKNRNLAFDVIK  
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repeat_region	15978..16009	/rpt_family="AT_rich"
repeat_region	comp1ement(16135..16202)	/rpt_family="(TA)n"
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Query Match	78.1%;	Score 16.4; DB 8; Length 68415;
Best Local Similarity	94.4%;	Pred. No. 7.3e+02;
Matches 17; Conservative	0; Mismatches	1; Indels 0; Gaps 0;
Oy	4 TCGAACGTTGCAGATGAT 21	
Db	23461 TCGATCGTTCCAGATGAT 23478	
RESULT 38		
AB024024/c		
LOCUS	AB024024	73921 bp DNA linear PLN 27-DEC-2000
DEFINITION	Arabidopsis thaliana genomic DNA, chromosome 5, TAC clone:K15C23.	
ACCESSION	AB024024 BA000015	
VERSION	AB024024.1 GI:4519183	
KEYWORDS		
SOURCE	Arabidopsis thaliana (thale cress)	

ORGANISM	REFERENCE AUTHORS	TITLE	JOURNAL	COMMENT
Arabidopsis thaliana	Eukaryote; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.	1 (stee)	Kaneke, T., Katoh, T., Asanlu, E., Sato, S., Nakamura, Y., Koreni, H. and Tabata, S.	Structural analysis of Arabidopsis thaliana chromosome 5. XI
Arabidopsis thaliana	Unpublished	2 (bases 1 to 73921)	Nakamura, Y.	Direct Submission
Arabidopsis thaliana	Submitted (24-FEB-1999)	Yasukazu Nakamura, Kazuo DNA Research Institute, Department of Plant Gene Research, 153-3, Yana, Kisarazu, Chiba 292-0812, Japan (E-mail: ynakamu@kazusa.or.jp, Tel:81-438-52-3935, Fax:81-438-52-3934)	Address for correspondence: kaso@kazusa.or.jp	For the latest information on annotation of this clone, please see <a href="http://www.kazusa.or.jp/kaos/cgi-bin/gsd_graph.cgi?c=K15C23">http://www.kazusa.or.jp/kaos/cgi-bin/gsd_graph.cgi?c=K15C23</a>
Arabidopsis thaliana	Genes with similarity to proteins in the databases are described in 'product' or 'note' qualifiers. Genes that have no significant protein similarity are described as 'unknown protein'.	The software programs used to predict genes include: Grail (Informatics Group, Oak Ridge National Laboratory, <a href="http://compio.ornl.gov/Grail-1.3/">http://compio.ornl.gov/Grail-1.3/</a> ), NetScan (Chris Burge, MIT, <a href="http://CCR-081.mit.edu/GENSCAN.html">http://CCR-081.mit.edu/GENSCAN.html</a> ), NetGene2 (S. M. Hebsgaard, et al., CBS, Technical University of Denmark, <a href="http://www.cbs.dtu.dk/services/NetGene2/">http://www.cbs.dtu.dk/services/NetGene2/</a> ) and SnpIcepredictor (Volker Brendel, Stanford University, <a href="http://gremlin.zool.iastate.edu/cgi-bin/sp.cgi">http://gremlin.zool.iastate.edu/cgi-bin/sp.cgi</a> )	Genes encoding tRNAs are predicted by tRNAscan-SE (Sean Eddy, Washington University School of Medicine, St. Louis, <a href="http://genome.wustl.edu/eddy/tRNAscan-SE/">http://genome.wustl.edu/eddy/tRNAscan-SE/</a> ).	This sequence may not be the entire insert of this clone. It may be shorter because we remove overlaps between neighboring submissions. The 5' clone is MRC16 and the 3' clone is K23L20.
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63301..64824

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GPAVYRPSISMSRIILPRGNLKGGINQAGIDYVNNLIELLSKGIKPEPFIITHMTPOS  
LEBAYGFGFSAEIVNDRDYADICFKNGDGRVKHMTNLEPLTVQOQYAVNAVGR  
CSFPTNKNCAANGATPEYIVGNHLIAHGSAVYVEEYKASQGVGILNAGMNL  
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unknown protein"

/codon\_start=1

/evidence=not\_experimental

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DLAEALQVNDGKIKPDRATLRMLHEMTIWPMLVEVSKRQKGSMAKSTGIDP  
KEAKRLNVEDSAAAIIEVDVDEQGVTVKAGYALYFVSALPVIIIGISVILIFY  
NSLD"

67965..69236

/note="gene\_id:K15C23.11

|||||T05327

similar to unknown protein"

/codon\_start=1

/evidence=not\_experimental

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/db\_xref="GI:8953764"

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REIMDPKSIIRLSMSGLSRNDSFDMRLPAMSPPRUDSPMLPLPVQTTGSPKO  
RSGMLRALNRDQSLPNSPKORSGLMVLAFFNKQDLSLPSNTTGSFKORSGMLRALNR  
KEQSLPNSNTGSPKORSGMLRALNRKQDSSASVYKSKSCGSTRKTLSHKSGSIRN

Query Match 78.1%; Score 16.4; DB 8; Length 73921;

Best Local Similarity 94.4%; Pred. No. 7.4e+02;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 TCGAACGTTTCGATGAT 21  
|||||

Db 38151 TCGAACGTTTCGATGAT 38134

RESULT 39

LOCUS AK100785/c 1687 bp mRNA linear PLN 24-JUL-2003

DEFINITION Oryza sativa (japonica cultivar-group) cDNA clone:022120N10, full  
insert sequence.

ACCESSION AK100785

VERSION AK100785.1 GI:32985994

KEYWORDS FLI\_CDNA; CAP\_trapper.

SOURCE Oryza sativa (japonica cultivar-group)

ORGANISM Oryza sativa (japonica cultivar-group); Embryophyta; Tracheophyta;  
Eukaryota; Viridiplantae; Streptophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
Ehhartrioideae; Oryzae; Oryza.

REFERENCE 1 The Rice Full-Length cDNA Consortium, National Institute of





RORDSIARIKRSRQTPNNVTYESRGGRRFFSRSERAGTVYSTIILPIRILIAN  
 TGLSTTSVAITDMLQJOWTGFELSYFMYSDADBPSPQWMDASBPQOGSS  
 SNNSTYSGSEGVFBSSNGSGTSGARREGRNTPGSSLPPLSLAQFLINE  
 DDDDDPRGLTJKOIDLINARFETALITCSYCTEYTEGNKLRILKPCSEHETHVCI  
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## ORIGIN

Query Match	77.1%	Score 16.2;	DB 5;	Length 2064;
Best Local Similarity	85.7%	Prod No. 6	3a402;	

Best Local Similarity 85.7%; Pred. No. 6.3e+02;  
Matches 18; Conservative 0; Mismatches 3

Matches	18;	Conservative	0;	Mismatches	3;	Indels	0;	Gaps	0;
---------	-----	--------------	----	------------	----	--------	----	------	----

QY 1 TCGTCGAACGTTGAGATGT 21

Db 894 TGGTCAACGTTTGAGATGAT 874

Search completed: April 6, 2004, 03:01:50  
Job time : 2885 secs

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: April 5, 2004, 21:30:52 ; Search time 396 Seconds  
(without alignments)  
225.283 Million cell updates/sec

Title: US-10-033-243-132

Perfect score: 21

Sequence: 1 tcgtcgacgttcgatgatg 21

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 3373863 seqs, 2124099041 residues

Total number of hits satisfying chosen parameters: 6747726

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database : N\_Geneseq\_29Jan04:\*

1: geneeqn1980s:\*\n2: geneeqn1990s:\*\n3: geneeqn2000s:\*\n4: geneeqn2001as:\*\n5: geneeqn2001bs:\*\n6: geneeqn2002s:\*\n7: geneeqn2003as:\*\n8: geneeqn2003bs:\*\n9: geneeqn2003cs:\*\n10: geneeqn2004s:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	21	100.0	21	6	ABQ75182 ISS immun
2	19	90.5	19	6	ABQ75170 ISS immun
3	19	90.5	19	7	ACC4937 Human imm
4	19	90.5	19	8	ADB88838 Chimeric
5	19	90.5	22	6	ABQ75181 ISS immun
6	19	90.5	22	8	ADB88849 Chimeric
7	18	85.7	19	6	ABQ75175 ISS immun
8	18	85.7	19	8	ADB88843 Chimeric
9	17.4	82.9	19	6	ABQ75221 ISS immun
10	17.4	82.9	19	6	ABQ75222 ISS immun
11	17.4	82.9	19	8	ADB88891 Chimeric
12	17.4	82.9	19	8	ADB88892 Chimeric
13	17	81.0	19	6	ABQ75174 ISS immun
14	17	81.0	19	8	ADB88842 Chimeric
15	16.4	78.1	18	8	ABQ75165 ISS immun
16	16.4	78.1	18	8	ADB88833 Chimeric
17	16.4	78.1	23	2	AAV80098 Immunomod
18	16.4	78.1	23	3	AAH38067 Immunosti
19	16.4	78.1	23	4	AAH75994 Immunomod
20	16.4	78.1	23	4	AAH77042 Immunosti
21	16.4	78.1	23	5	AAH41575 Immunosti
22	16.4	78.1	23	5	AAH14666 Immunosti
23	16.4	78.1	23	6	ABA03835 Immunosti

C	24	16.4	78.1	23	6	ABA03846	ABa03846 Immunosti
C	25	16.4	78.1	23	6	AAH36339	AAH36339 ISS polyn
C	26	16.4	78.1	23	6	AAH36350	AAH36350 ISS polyn
C	27	16.4	78.1	23	6	ABA03858	ABA03858 Immunosti
C	28	16.2	77.1	1017	4	ABL12591	ABL12591 Drosophi
C	29	16.2	77.1	1020	7	ACA47671	ACA47671 Prokaryot
C	30	16.2	77.1	1676	6	ABQ14749	ABQ14749 Oligonuc
C	31	16.2	77.1	1676	6	ABQ14748	ABQ14748 Oligonuc
C	32	16.2	77.1	3017	4	ABL12594	ABL12594 Drosophi
C	33	16.2	77.1	3126	7	ACA33960	ACA33960 Prokaryot
C	34	16.2	77.1	3135	9	ADC91948	ADC91948 E. faeciu
C	35	16.2	77.1	4024	2	AAV17086	AAV17086 Bos tauru
C	36	16	76.2	66	8	ADB88936	ADB88936 Chimeric
C	37	15.8	75.2	19	8	ABQ75223	ABQ75223 ISS immun
C	38	15.8	75.2	19	8	ADB88893	ADB88893 Chimeric
C	39	15.8	75.2	2726	4	ABL09441	ABL09441 Drosophi
C	40	15.8	75.2	5080	4	ABL09440	ABL09440 Drosophi
C	41	15.8	75.2	6128	6	ABQ67039	ABQ67039 Human ang
C	42	15.8	75.2	6129	6	ABK31236	ABK31236 Signal tr
C	43	15.8	75.2	6129	6	ABL70537	ABL70537 Chemical
C	44	15.8	75.2	6129	6	AAH61149	AAH61149 Human gen
C	45	15.8	75.2	11131	4	ABL09428	ABL09428 Drosophi

## ALIGNMENTS

RESULT 1	ABQ75182 standard; DNA; 21 BP.
ID	ABQ75182
XX	XX
AC	ABQ75182;
XX	XX
DT	05-NOV-2002 (first entry)
XX	XX
DE	ISS immunomodulatory oligonucleotide SEQ ID NO:132.
XX	XX
KW	Immunostimulatory sequence; ISS: immunomodulatory; immune response; allergy; asthma; infectious disease; interferon-gamma; IFN-gamma; idiopathic pulmonary fibrosis; infection; mycobacterial disease; malaria; leishmaniasis; toxoplasmosis; schistosomiasis; immunoglobulin E; IGE-related disorder; antiallergic; antiasthmatic; vituicide; antibacterial; protozoacide; ss.
KW	XX
OS	Synthetic.
XX	XX
PN	WO200252002-A2.
XX	XX
PD	04-JUL-2002.
XX	XX
PF	27-DEC-2001; 2001WO-US050821.
XX	XX
PR	27-DEC-2000; 2000US-0258675P.
XX	XX
PA	(DYNA-) DYNAVAX TECHNOLOGIES CORP.
XX	XX
P1	Fearon KL, Dina D;
XX	XX
DR	WPI: 2002-657426/70.
XX	XX
PT	Immunomodulatory polynucleotide for modulating an immune response in a subject suffering from disorders associated with Th2-type immune response, e.g. allergy, or infectious disease, comprises an immunostimulatory sequence.
PT	XX
PS	Claim 4; Page 21; 95pp; English.
XX	XX
XX	XX
CC	The present invention describes an immunomodulatory polynucleotide (1) comprising an immunostimulatory sequence (ISS). Also described: (1) an immunomodulatory composition comprising (1); (2) an immunomodulatory polynucleotide/microcarrier (IMP/MC) complex, comprising (1) linked to a biodegradable MC, where the MC is less than 10 micrometre in size; and (3) a kit comprising (1). (1) has antiallergic, antiasthmatic, vituicide,
CC	CC

CC antibacterial and protozoacide activities, and can be used as a modulator  
CC of immune response. (1) is useful for modulating an immune response in an  
CC individual suffering from disorders associated with a Th2-type immune  
CC response, especially an allergy or asthma, or an infectious disease. (1)  
CC is also useful for increasing interferon-gamma (IFN-gamma) in an  
CC individual having idiopathic pulmonary fibrosis, or IFN-alpha in an  
CC individual having a viral infection. (1) is further useful for  
CC ameliorating a symptom of an infectious disease caused by a cellular  
CC pathogen such as mycobacterial disease, malaria, leishmaniasis,  
CC toxoplasmosis, schistosomiasis and clonorchiasis in an individual, or a  
CC symptom of an immunoglobulin E (IGE)-related disorder, preferably an  
CC allergy-related disorder, in particular asthma in an individual. The  
CC present sequence represents an immunomodulatory oligonucleotide from the  
CC present invention

XX  
XX Sequence 21 BP; 5 A; 4 C; 6 G; 6 T; 0 U; 0 Other;  
SQ

Query Match 100.0%; Score 21; DB 6; Length 21;  
Best Local Similarity 100.0%; Pred. No. 0.26;  
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCGTGAACGTTGAGATGAT 21  
1 TCGTGAACGTTGAGATGAT 21

DB 1 TCGTGAACGTTGAGATGAT 21

RESULT 2  
ABQ75170  
ID ABQ75170 standard; DNA; 19 BP;  
XX  
XX ABQ75170;  
AC  
XX  
XX 05-NOV-2002 (first entry)  
DT

ISS immunomodulatory oligonucleotide SEQ ID NO:19.  
XX  
XX Immunostimulatory sequence; ISS: immunomodulatory; immune response;  
XX allergy; asthma; infectious disease; interferon-gamma; IFN-gamma;  
XX idiopathic pulmonary fibrosis; viral infection; mycobacterial disease;  
XX malaria; leishmaniasis; toxoplasmosis; schistosomiasis; clonorchiasis;  
XX immunoglobulin E; IGE-related disorder; antiallergic; antiasthmatic;  
XX virucide; antibacterial; protozoacide; ss.  
XX  
XX Synthetic.  
OS  
XX  
XX WO200252002-A2.  
PN  
XX  
XX 04-JUL-2002.  
PD  
XX  
XX 27-DEC-2001; 2001WO-US050821.  
PF  
XX  
XX 27-DEC-2000; 2000US-0258675P.  
PR  
XX  
XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
PA  
XX  
XX Fearon KL, Dina D;  
PI  
XX  
XX WPI; 2002-657426/70.  
DR

Immunomodulatory polynucleotide for modulating an immune response in a  
PT subject suffering from disorders associated with Th2-type immune  
PT response, e.g. allergy, or infectious disease, comprises an  
PT immunostimulatory sequence.  
XX  
XX  
XX Claim 4; Page 20; 95pp; English.  
PS

The present invention describes an immunomodulatory polynucleotide (1)  
CC comprising an immunostimulatory sequence (ISS). Also described: (1) an  
CC immunomodulatory composition comprising (1); (2) an immunomodulatory  
CC polynucleotide/microcarrier (IMP/MC) complex, comprising (1) linked to a  
CC biodegradable MC, where the MC is less than 10 micrometre in size; and  
CC (3) a kit comprising (1). (1) has antiallergic, antiasthmatic, virucide,  
CC antibacterial and protozoacide activities, and can be used as a modulator

CC of immune response. (1) is useful for modulating an immune response in an  
CC individual suffering from disorders associated with a Th2-type immune  
CC response, especially an allergy or asthma, or an infectious disease. (1)  
CC is also useful for increasing interferon-gamma (IFN-gamma) in an  
CC individual having idiopathic pulmonary fibrosis, or IFN-alpha in an  
CC individual having a viral infection. (1) is further useful for  
CC ameliorating a symptom of an infectious disease caused by a cellular  
CC pathogen such as mycobacterial disease, malaria, leishmaniasis,  
CC toxoplasmosis, schistosomiasis and clonorchiasis in an individual, or a  
CC symptom of an immunoglobulin E (IGE)-related disorder, preferably an  
CC allergy-related disorder, in particular asthma in an individual. The  
CC present sequence represents an immunomodulatory oligonucleotide from the  
CC present invention

XX  
XX Sequence 19 BP; 4 A; 4 C; 6 G; 5 T; 0 U; 0 Other;  
SQ

Query Match 90.5%; Score 19; DB 6; Length 19;  
Best Local Similarity 100.0%; Pred. No. 3;  
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCGTGAACGTTGAGATG 19  
1 TCGTGAACGTTGAGATG 19

DB 1 TCGTGAACGTTGAGATG 19

RESULT 3  
ACC49937  
ID ACC49937 standard; DNA; 19 BP.  
XX  
XX ACC49937;  
AC  
XX  
XX 26-JUN-2003 (first entry)  
DT

Human immunomodulatory polynucleotide SEQ ID NO: 11.  
XX  
XX Human immunomodulatory polynucleotide; microcarrier complex;  
XX Human; IMP; immunomodulatory; antiinflammatory; cytostatic; virucide;  
XX antiasthmatic; antiallergic; antitumor; dermatological; immune response;  
XX hepatotropic; protozoacide; antitumor; dermatological; immune response;  
XX interferon-gamma; interferon-alpha; IGEF; vaccine; asthma; rhinitis;  
XX allergic conjunctivitis; cancer; infectious disease; viral infection;  
XX parasitic infection; inflammatory disorder; ss.  
XX  
XX Homo sapiens.  
OS  
XX  
XX WO2003015816-A1.  
PN  
XX  
XX 27-FEB-2003.  
PD  
XX  
XX 13-AUG-2001; 2001WO-US025364.  
PF  
XX  
XX 10-AUG-2001; 2001US-00927422.  
PR  
XX  
XX 10-AUG-2001; 2001US-00927884.  
PA  
XX  
XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
PA  
XX  
XX Van Nest G, Tuck S, Fearon KL, Dina D;  
PI  
XX  
XX WPI; 2003-312719/30.  
DR

Immunomodulatory polynucleotide/microcarrier complexes that can increase  
PT levels of interferon-gamma and interferon-alpha and reduce levels of  
PT IGEF, comprises heptameric oligonucleotides.  
XX  
XX  
XX Example 5; Page 61; 84pp; English.  
PS

The invention relates to novel immunomodulatory polynucleotide/  
CC microcarrier complexes comprising heptameric oligonucleotides comprising  
CC a 5' CG-3' sequence. A complex of the invention has antiasthmatic,  
CC antiallergic, antiinflammatory, cytostatic, virucide, hepatotropic,  
CC protozoacide, antitumor, and dermatological activity. The complexes are  
CC useful for modulating an immune response, increasing interferon-gamma,  
CC increasing interferon-alpha, and reducing levels of IGEF. The complexes  
CC can also be used in vaccines, to treat, for example, asthma, rhinitis,

CC allergic conjunctivitis, cancer, infectious diseases, viral infections,  
CC parasitic infections (e.g. malaria, leishmaniasis), inflammatory  
CC disorders (e.g. ulcerative colitis, idiopathic pulmonary fibrosis,  
CC scleroderma). The present sequence is used in the exemplification of the  
CC invention.

XX SQ Sequence 19 BP; 4 A; 4 C; 6 G; 5 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 7; Length 19;  
Best Local Similarity 100.0%; Pred. No. 3;  
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGAACGTTCCGAGATG 19  
DB 1 TCGTCGAACGTTCCGAGATG 19

## RESULT 4

ADB88838  
ID ADB88838 standard; DNA; 19 BP.

XX ADB88838;

XX 04-DEC-2003 (first entry)

XX Chimeric immunomodulatory compound DNA sequence, SEQ ID No 41.

XX chimeric immunomodulatory compound; CIC; immunomodulatory activity;

XX spacer moiety; linear hexaethylene glycol structure; HEG; immune;

XX Th2-type; allergy; allergy-induced asthma; infectious disease; IFN-gamma;

XX IFN-alpha; idiopathic pulmonary fibrosis; viral infection; ameliorating;

XX immunoglobulin E; IgE; allergy; cancer;

XX stimulating cellular immune system cell; ss.

XX Synthetic.

XX WO200300922-A2.

XX 03-JAN-2003.

XX 21-JUN-2002; 2002WO-US020025.

XX 21-JUN-2001; 2001US-0299883P.

XX 23-APR-2002; 2002US-0375253P.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX Fearon KL, Dina D, Tuck SF;

XX WPI; 2003-210159/20.

XX The invention relates to a novel chimeric immunomodulatory compound (CIC)

XX having immunomodulatory activity, comprising two or more nucleic acid

XX moieties and one or more non-nucleic acid spacer moieties, where at least

XX one non-nucleic acid spacer moiety is covalently joined to two nucleic

XX acid moieties, where the spacer is not a polypeptide, and at least one

XX nucleic acid moiety comprises the sequence 5'-CG-3'. The chimeric

XX immunomodulatory compound more specifically contain the nucleic acid

XX spacer moieties of linear hexaethylene glycol structure (HEG) subunits.

XX CIC's are useful for modulating an immune response in an individual,

XX where the individual suffers from a disorder associated with a Th2-type

XX immune response which is an allergy or allergy-induced asthma, and an

XX infectious disease. CIC is also useful for increasing IFN-gamma, or IFN-

XX alpha, in an individual, where the individual has idiopathic pulmonary

XX fibrosis, or a viral infection. CIC's are useful for ameliorating a

XX symptom of an infectious disease, or an immunoglobulin E (IgE)-related

XX disorder in an individual, where the IgE-related disorder is allergy, or

CC an allergy-related disorder. CIC's are also useful for treating cancer

CC and can be used for stimulating cellular immune system cells production

CC in an individual. This polynucleotide sequence represents a DNA sequence

CC which is a nucleic acid moiety part of a chimeric immunomodulatory compound

CC of the invention.

XX SQ Sequence 19 BP; 4 A; 4 C; 6 G; 5 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 8; Length 19;  
Best Local Similarity 100.0%; Pred. No. 3;  
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGAACGTTCCGAGATG 19  
DB 1 TCGTCGAACGTTCCGAGATG 19

## RESULT 5

ABQ75181  
ID ABQ75181 standard; DNA; 22 BP.

XX ABQ75181;

XX 05-NOV-2002 (first entry)

XX ISS immunomodulatory oligonucleotide SEQ ID NO:30.

XX immunostimulatory sequence; ISS: immunomodulatory; immune response;

XX allergy; asthma; infectious disease; interferon-gamma; IFN-gamma;

XX idiopathic pulmonary fibrosis; viral infection; mycobacterial disease;

XX malaria; leishmaniasis; toxoplasmosis; schistosomiasis; clonorchiasis;

XX immunoglobulin E; IgE-related disorder; antiallergic; antiasthmatic;

XX virucide; antibacterial; protozoacide; ss.

XX Synthetic.

XX WO200252002-A2.

XX 04-JUL-2002.

XX 27-DEC-2001; 2001WO-US050821.

XX 27-DEC-2000; 2000US-0258675P.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX Fearon KL, Dina D;

XX WPI; 2002-657426/70.

XX The present invention describes an immunomodulatory polynucleotide (I)

XX comprising an immunostimulatory sequence (ISS). Also described: (1) an

XX immunomodulatory composition comprising (I); (2) an immunomodulatory

XX polynucleotide/microcarrier (IMP/MC) complex, comprising (I) linked to a

XX biodegradable MC, where the MC is less than 10 micrometre in size, and

XX (3) a kit comprising (1). (1) has antiallergic, antiasthmatic, virucide,

XX antibacterial and protozoacide activities, and can be used as a modulator

XX of immune response. (I) is useful for modulating an immune response in an

XX individual suffering from disorders associated with a Th2-type immune

XX response, especially an allergy or asthma, or an infectious disease. (I)

XX is also useful for increasing interferon-gamma (IFN-gamma) in an

XX individual having idiopathic pulmonary fibrosis, or IFN-alpha in an

XX individual having a viral infection. (I) is further useful for

XX ameliorating a symptom of an infectious disease caused by a cellular

XX pathogen such as mycobacterial disease, malaria, leishmaniasis,

XX toxoplasmosis, schistosomiasis and clonorchiasis in an individual, or a

CC symptom of an immunoglobulin E (IGE)-related disorder, preferably an  
 CC allergy-related disorder, in particular asthma in an individual. The  
 CC present sequence represents an immunomodulatory oligonucleotide from the  
 CC present invention

XX Sequence 22 BP; 5 A; 4 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 6; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 3;  
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTGAACGTTGAGATG 19  
 |||||  
 DB 4 TCGTGAACGTTGAGATG 22

RESULT 6  
 ADB88849  
 ID ADB88849 standard; DNA; 22 BP.

XX ADB88849;

DT 04-DEC-2003 (first entry)

XX Chimeric immunomodulatory compound DNA sequence, SEQ ID No 52.

DE Chimeric immunomodulatory compound DNA sequence, SEQ ID No 52.  
 XX chimeric immunomodulatory compound; CTC; immunomodulatory activity;  
 KW spacer moiety; linear hexaethylene glycol structure; HEG; immune;  
 KW Th2-type; allergy; allergy-induced asthma; infectious disease; IFN-gamma;  
 KW IFN-alpha; idiopathic pulmonary fibrosis; viral infection; ameliorating;  
 KW immunoglobulin E; IGE; allergy; cancer;  
 KW stimulating cellular immune system cell; ss.

XX Synthetic.

OS WO200300922-A2.

XX 03-JAN-2003.

PD 21-JUN-2002; 2002WO-US020025.

XX 21-JUN-2001; 2001US-0299883P.

PR 23-APR-2002; 2002US-0375253P.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

PA Fearon KL, Dina D, Tuck SF;

PI WPI; 2003-210159/20.

PT Novel chimeric immunomodulatory compound having immunomodulatory  
 PT activity, useful for modulating an immune response and for treating  
 PT cancer, has nucleic acid moieties and non-nucleic acid spacer moieties.

XX Disclosure; Page 34; 224pp; English.

CC The invention relates to a novel chimeric immunomodulatory compound (CIC)  
 CC having immunomodulatory activity, comprising two or more nucleic acid  
 CC moieties and one or more non-nucleic acid spacer moieties, where at least  
 CC one non-nucleic acid spacer moiety is covalently joined to two nucleic  
 CC acid moieties, where the spacer is not a polypeptide, and at least one  
 CC nucleic acid moiety comprises the sequence 5'-CG-3'. The chimeric  
 CC immunomodulatory compounds more specifically contain the nucleic acid  
 CC spacer moieties of linear hexaethylene glycol structure (HEG) subunits.  
 CC CIC's are useful for modulating an immune response in an individual,  
 CC where the individual suffers from a disorder associated with a Th2-type  
 CC immune response which is an allergy or allergy-induced asthma, and an  
 CC infectious disease. CIC is also useful for increasing IFN-gamma; or IFN-  
 CC alpha; in an individual, where the individual has idiopathic pulmonary  
 CC fibrosis, or a viral infection. CIC's are useful for ameliorating a  
 CC symptom of an infectious disease, or an immunoglobulin E (IGE)-related  
 CC disorder in an individual, where the IGE-related disorder is allergy, or  
 CC an allergy-related disorder. CIC's are also useful for treating cancer

CC and can be used for stimulating cellular immune system cells production  
 CC in an individual. This polynucleotide sequence represents a DNA sequence  
 CC which is a nucleic acid moiety part of a chimeric immunomodulatory compound  
 CC of the invention.

XX Sequence 22 BP; 5 A; 4 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 8; Length 22;  
 Best Local Similarity 100.0%; Pred. No. 3;  
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTGAACGTTGAGATG 19  
 |||||  
 DB 4 TCGTGAACGTTGAGATG 22

RESULT 7  
 ABQ75175  
 ID ABQ75175 standard; DNA; 19 BP.

XX ABQ75175;

DT 05-NOV-2002 (first entry)

XX ISS immunomodulatory oligonucleotide SEQ ID NO:24.

DE Immunostimulatory sequence; ISS; immunomodulatory; immune response;  
 KW allergy; asthma; infectious disease; interferon-gamma; IFN-gamma;  
 KW idiopathic pulmonary fibrosis; viral infection; mycobacterial disease;  
 KW malaria; leishmaniasis; toxoplasmosis; schistosomiasis; clonorchiasis;  
 KW immunoglobulin E; IGE-related disorder; antiallergic; antiasthmatic;  
 KW virucide; antibacterial; protozoacide; ss.

XX Synthetic.

OS Key Location/Qualifiers

XX modified\_base 5

FT /\*tag= a

FT /mod\_base= OTHER

FT /note="5-bromocytosine"

XX WO200252002-A2.

XX 04-JUL-2002.

PD 27-DEC-2001; 2001WO-US050821.

XX 27-DEC-2000; 2000US-0258675P.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

PA Fearon KL, Dina D;

PI WPI; 2002-657426/70.

PT Immunomodulatory polynucleotide for modulating an immune response in a  
 PT subject suffering from disorders associated with Th2-type immune  
 PT response, e.g. allergy, or infectious disease, comprises an  
 PT immunostimulatory sequence.

XX Example 1; Page 20; 95pp; English.

CC The present invention describes an immunomodulatory polynucleotide (I)  
 CC comprising an immunostimulatory sequence (ISS). Also described: (1) an  
 CC immunomodulatory composition comprising (I); (2) an immunomodulatory  
 CC polynucleotide/microcarrier (IMP/MC) complex, comprising (I) linked to a  
 CC biodegradable MC, where the MC is less than 10 micrometre in size; and  
 CC (3) a kit comprising (I). (I) has antiallergic, antiasthmatic, virucide,  
 CC antibacterial and protozoacide activities, and can be used as a modulator  
 CC of immune response. (I) is useful for modulating an immune response in an  
 CC individual suffering from disorders associated with a Th2-type immune  
 CC response, especially an allergy or asthma, or an infectious disease. (I)  
 CC is also useful for increasing interferon-gamma (IFN-gamma) in an

CC individual having idiopathic pulmonary fibrosis, or IFN-alpha in an  
CC individual having a viral infection. (1) is further useful for  
CC ameliorating a symptom of an infectious disease caused by a cellular  
CC pathogen such as mycobacterial disease, malaria, leishmaniasis,  
CC toxoplasmosis, schistosomiasis and clonorchiasis in an individual, or a  
CC symptom of an immunoglobulin E (IGE)-related disorder, preferably an  
CC allergy-related disorder, in particular asthma in an individual. The  
CC present sequence represents an immunomodulatory oligonucleotide from the  
CC present invention.

XX Sequence 19 BP; 4 A; 3 C; 6 G; 5 T; 0 U; 1 Other;

Query Match 85.7%; Score 18; DB 6; Length 19;  
Best Local Similarity 94.7%; Pred. No. 10;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 TCGTGAACGTTCCGAGATG 19  
Db 1 TCGTNGAACGTTCCGAGATG 19

RESULT 8  
ADB88843

ID ADB88843 standard; DNA; 19 BP.

XX ADB88843;

DT 04-DEC-2003 (first entry)

XX Chimeric immunomodulatory compound DNA sequence, SEQ ID No 46.

KM chimeric immunomodulatory compound; CIC; immunomodulatory activity;  
KM spacer moiety; linear hexethylene glycol structure; HEG; immune;  
KM Th2-type; allergy; allergy-induced asthma; infectious disease; IFN-gamma;  
KM IFN-alpha; idiopathic pulmonary fibrosis; viral infection; ameliorating;  
KM immunoglobulin E; IGE; allergy; cancer;  
KM stimulating cellular immune system cell; ss.

XX Synthetic.

OS WO2003000922-A2.

PN 03-JAN-2003.

PD 21-JUN-2002; 2002WO-US020025.

XX 21-JUN-2001; 2001US-029883P.

PR 23-APR-2002; 2002US-0375253P.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

PA Fearon KL, Dina D, Tuck SF;

PI WPI; 2003-210159/20.

XX Novel chimeric immunomodulatory compound having immunomodulatory  
PT activity, useful for modulating an immune response and for treating  
PT cancer, has nucleic acid moieties and non-nucleic acid spacer moieties.

XX Disclosure; Page 33; 224pp; English.

PS The invention relates to a novel chimeric immunomodulatory compound (CIC)  
CC having immunomodulatory activity, comprising two or more nucleic acid  
CC moieties and one or more non-nucleic acid spacer moieties, where at least  
CC one non-nucleic acid spacer moiety is covalently joined to two nucleic  
CC acid moieties, where the spacer is not a polypeptide, and at least one  
CC nucleic acid moiety comprises the sequence 5'-GG-3'. The chimeric  
CC immunomodulatory compound more specifically contain the nucleic acid  
CC spacer moieties of linear hexethylene glycol structure (HEG) subunits.  
CC CIC's are useful for modulating an immune response in an individual,  
CC where the individual suffers from a disorder associated with a Th2-type  
CC immune response which is an allergy or allergy-induced asthma, and an  
CC infectious disease. CIC is also useful for increasing IFN-gamma, or IFN-

CC alpha; in an individual, where the individual has idiopathic pulmonary  
CC fibrosis, or a viral infection. CIC's are useful for ameliorating a  
CC symptom of an infectious disease, or an immunoglobulin E (IGE)-related  
CC disorder in an individual, where the IGE-related disorder is allergy, or  
CC an allergy-related disorder. CIC's are also useful for treating cancer  
CC and can be used for stimulating cellular immune system cells production  
CC in an individual. This polynucleotide sequence represents a DNA sequence  
CC which is a nucleic acid moiety part of a chimeric immunomodulatory compound  
CC of the invention.

XX Sequence 19 BP; 4 A; 3 C; 6 G; 5 T; 0 U; 1 Other;

Query Match 85.7%; Score 18; DB 8; Length 19;  
Best Local Similarity 94.7%; Pred. No. 10;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 TCGTGAACGTTCCGAGATG 19  
Db 1 TCGTNGAACGTTCCGAGATG 19

RESULT 9  
ABQ75221

ID ABQ75221 standard; DNA; 19 BP.

XX ABQ75221;

DT 05-NOV-2002 (first entry)

XX ISS immunomodulatory oligonucleotide SEQ ID NO:56.

KM Immunostimulatory sequence; ISS; immunomodulatory; immune response;  
KM allergy; asthma; infectious disease; interferon-gamma; IFN-gamma;  
KM idiopathic pulmonary fibrosis; viral infection; mycobacterial disease;  
KM malaria; leishmaniasis; toxoplasmosis; schistosomiasis; clonorchiasis;  
KM immunoglobulin E; IGE-related disorder; antiallergic; antiasthmatic;  
KM virucide; antibacterial; protozoacide; ss.

XX Synthetic.

OS WO200252002-A2.

PN 04-JUL-2002.

PD 27-DEC-2001; 2001WO-US050821.

XX 27-DEC-2000; 2000US-0258675P.

PR (DYNA-) DYNAVAX TECHNOLOGIES CORP.

PA Fearon KL, Dina D;

PI WPI; 2002-657426/70.

XX Immunomodulatory polynucleotide for modulating an immune response in a  
PT subject suffering from disorders associated with Th2-type immune  
PT response, e.g. allergy, or infectious disease, comprises an  
PT immunostimulatory sequence.

XX Disclosure; Page 22; 95pp; English.

PS The present invention describes an immunomodulatory polynucleotide (1)  
CC comprising an immunostimulatory sequence (ISS). Also described: (1) an  
CC immunomodulatory composition comprising (1); (2) an immunomodulatory  
CC polynucleotide/microcarrier (IMP/MC) complex, comprising (1) linked to a  
CC biodegradable MC, where the MC is less than 10 micrometre in size; and  
CC (3) a kit comprising (1). (1) has antiallergic, antiasthmatic, virucide,  
CC antibacterial and protozoacide activities, and can be used as a modulator  
CC of immune response. (1) is useful for modulating an immune response in an  
CC individual suffering from disorders associated with a Th2-type immune  
CC response, especially an allergy or asthma, or an infectious disease. (1)  
CC is also useful for increasing interferon-gamma (IFN-gamma) in an  
CC individual having idiopathic pulmonary fibrosis, or IFN-alpha in an





CC symptom of an infectious disease, or an immunoglobulin E (IgE)-related  
CC disorder in an individual, where the IgE-related disorder is allergy, or  
CC an allergy-related disorder. CIC's are also useful for treating cancer  
CC and can be used for stimulating cellular immune system cells production  
CC in an individual. This polynucleotide sequence represents a DNA sequence  
CC which is a nucleic acid moiety part of a chimeric immunomodulatory compound  
CC of the invention.

XX  
SQ Sequence 19 BP; 3 A; 4 C; 7 G; 5 T; 0 U; 0 Other;

Query Match 82.9%; Score 17.4; DB 8; Length 19;  
Best Local Similarity 94.7%; Pred. No. 21;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 TCGTCGAACGTTGAGATG 19  
DB 1 TCGTCGACGTTGAGATG 19  
|||||  
|||||

RESULT 12  
ADB88892  
ID ADB88892 standard; DNA; 19 BP.  
XX  
AC ADB88892;  
XX  
DT 04-DEC-2003 (first entry)  
XX  
DE Chimeric immunomodulatory compound DNA sequence, SEQ ID NO 95.  
XX  
XX chimeric immunomodulatory compound; CIC; immunomodulatory activity;  
KW spacer moiety; linear hexaethylene glycol structure; HEG; immune;  
KW Th2-type; allergy; allergy-induced asthma; infectious disease; IFN-gamma;  
KW IFN-alpha; idiopathic pulmonary fibrosis; viral infection; ameliorating;  
KW immunoglobulin E; IgE; allergy; cancer;  
KW stimulating cellular immune system cell; ss.  
XX  
OS Synthetic.  
XX  
PN WO200300922-A2.  
XX  
PD 03-JAN-2003.  
XX  
PF 21-JUN-2002; 2002MO-US020025.  
XX  
PR 21-JUN-2001; 2001US-0299883P.  
XX  
PR 23-APR-2002; 2002US-0375253P.  
XX  
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
XX  
PI Fearon KL, Dina D, Tuck SF;  
XX  
DR WPI, 2003-210159/20.  
XX  
PT Novel chimeric immunomodulatory compound having immunomodulatory  
PT activity, useful for modulating an immune response and for treating  
PT cancer, has nucleic acid moieties and non-nucleic acid spacer moieties.  
XX  
PS Disclosure; Page 35; 224pp; English.  
XX  
CC The invention relates to a novel chimeric immunomodulatory compound (CIC)  
CC having immunomodulatory activity, comprising two or more nucleic acid  
CC moieties and one or more non-nucleic acid spacer moieties, where at least  
CC one non-nucleic acid spacer moiety is covalently joined to two nucleic  
CC acid moieties, where the spacer is not a polypeptide, and at least one  
CC nucleic acid moiety comprises the sequence 5'-CG-3'. The chimeric  
CC immunomodulatory compound more specifically contain the nucleic acid  
CC spacer moieties of linear hexaethylene glycol structure (HEG) subunits.  
CC CIC's are useful for modulating an immune response in an individual,  
CC where the individual suffers from a disorder associated with a Th2-type  
CC immune response which is an allergy or allergy-induced asthma, and an  
CC infectious disease. CIC is also useful for increasing IFN-gamma, or IFN-  
CC alpha, in an individual, where the individual has idiopathic pulmonary  
CC fibrosis, or a viral infection. CIC's are useful for ameliorating a

CC symptom of an infectious disease, or an immunoglobulin E (IgE)-related  
CC disorder in an individual, where the IgE-related disorder is allergy, or  
CC an allergy-related disorder. CIC's are also useful for treating cancer  
CC and can be used for stimulating cellular immune system cells production  
CC in an individual. This polynucleotide sequence represents a DNA sequence  
CC which is a nucleic acid moiety part of a chimeric immunomodulatory compound  
CC of the invention.

XX  
SQ Sequence 19 BP; 3 A; 4 C; 6 G; 6 T; 0 U; 0 Other;

Query Match 82.9%; Score 17.4; DB 8; Length 19;  
Best Local Similarity 94.7%; Pred. No. 21;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 TCGTCGAACGTTGAGATG 19  
DB 1 TCGTCGACGTTGAGATG 19  
|||||  
|||||

RESULT 13  
ABQ75174  
ID ABQ75174 standard; DNA; 19 BP.  
XX  
AC ABQ75174;  
XX  
DT 05-NOV-2002 (first entry)  
XX  
DE ISS immunomodulatory oligonucleotide SEQ ID NO:23.  
XX  
XX Immunostimulatory sequence; ISS; immunomodulatory; immune response;  
KW allergy; asthma; infectious disease; interferon-gamma; IFN-gamma;  
KW idiopathic pulmonary fibrosis; viral infection; mycobacterial disease;  
KW malaria; leishmaniasis; toxoplasmosis; schistosomiasis; clonorchiasis;  
KW immunoglobulin E; IgE-related disorder; anti-allergic; antiasthmatic;  
KW virucide; antibacterial; protozoacide; ss.  
XX  
OS Synthetic.  
XX  
PN WO200252002-A2.  
XX  
PD 04-JUL-2002.  
XX  
PF 27-DEC-2001; 2001WO-US050821.  
XX  
PR 27-DEC-2000; 2000US-0258675P.  
XX  
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
XX  
PI Fearon KL, Dina D;  
XX  
DR WPI, 2002-657426/70.  
XX  
PT Immunomodulatory polynucleotide for modulating an immune response in a  
PT subject suffering from disorders associated with Th2-type immune  
PT response, e.g. allergy, or infectious disease, comprises an  
PT immunostimulatory sequence.  
XX  
PS Example 1; Page 20; 95pp; English.  
XX  
CC The present invention describes an immunomodulatory polynucleotide (I)  
CC comprising an immunostimulatory sequence (ISS). Also described: (1) an  
CC immunomodulatory composition comprising (i); (2) an immunomodulatory  
CC polynucleotide/microcarrier (IMP/MC) complex, comprising (I) linked to a

CC biodegradable MC, where the MC is less than 10 micrometre in size; and  
CC (3) a kit comprising (1), (1) has anti-allergic, antiaesthetic, virucide,  
CC antibacterial and protozoacide activities, and can be used as a modulator  
CC of immune response. (1) is useful for modulating an immune response in an  
CC individual suffering from disorders associated with a Th2-type immune  
CC response, especially an allergy or asthma, or an infectious disease. (1)  
CC is also useful for increasing interferon-gamma (IFN-gamma) in an  
CC individual having idiopathic pulmonary fibrosis, or IFN-alpha in an  
CC individual having a viral infection. (1) is further useful for  
CC ameliorating a symptom of an infectious disease caused by a cellular  
CC pathogen such as mycobacterial disease, malaria, leishmaniasis,  
CC toxoplasmosis, schistosomiasis and clonorchiasis in an individual, or a  
CC symptom of an immunoglobulin E (IgE)-related disorder, preferably an  
CC allergy-related disorder, in particular asthma in an individual. The  
CC present sequence represents an immunomodulatory oligonucleotide from the  
CC present invention

XX  
SQ Sequence 19 BP; 4 A; 2 C; 6 G; 5 T; 0 U; 2 Other;

Query Match 81.0%; Score 17; DB 6; Length 19;  
Best Local Similarity 89.5%; Pred. No. 34;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGACGTTGAGATG 19  
| | | | | | | | | | | | | | | | | | |  
Db 1 TNGTNGACGTTGAGATG 19

## RESULT 14

ADB88842  
ID ADB88842 standard; DNA; 19 BP.

XX  
AC ADB88842;

DT 04-DEC-2003 (first entry)

XX  
DE Chimeric immunomodulatory compound DNA sequence, SEQ ID No 45.

XX  
KM chimeric immunomodulatory compound; C1C; immunomodulatory activity;  
KM spacer moiety; linear hexaethylene glycol structure; HEG; immune;  
KM Th2-type; allergic; allergy-induced asthma; infectious disease; IFN-gamma;  
KM IFN-alpha; idiopathic pulmonary fibrosis; viral infection; ameliorating;  
KM immunoglobulin E; IgE; allergy; cancer;  
KM stimulating cellular immune system cell; ss.

XX  
OS Synthetic.

XX  
PN WO2003000922-A2.

XX  
PD 03-JAN-2003.

XX  
PF 21-JUN-2002; 2002WO-US020025.

XX  
PR 21-JUN-2001; 2001US-0299883P.

XX  
PR 23-APR-2002; 2002US-0375253P.

XX  
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX  
PI Fearon KL, Dina D, Tuck SF;

XX  
DR WPI; 2003-210159/20.

XX  
PT Novel chimeric immunomodulatory compound having immunomodulatory  
PT activity, useful for modulating an immune response and for treating  
PT cancer, has nucleic acid moieties and non-nucleic acid spacer moieties.  
XX  
PS Disclosure; Page 33; 224pp; English.

CC The invention relates to a novel chimeric immunomodulatory compound (C1C)  
CC comprising an immunomodulatory activity, comprising two or more nucleic acid  
CC moieties and one or more non-nucleic acid spacer moieties, where at least  
CC one non-nucleic acid spacer moiety is covalently joined to two nucleic  
CC acid moieties, where the spacer is not a polypeptide, and at least one

CC nucleic acid moiety comprises the sequence 5'-CG-3'. The chimeric  
CC immunomodulatory compounds more specifically contain the nucleic acid  
CC spacer moieties of linear hexaethylene glycol structure (HEG) subunits.  
CC C1C's are useful for modulating an immune response in an individual,  
CC where the individual suffers from a disorder associated with a Th2-type  
CC immune response which is an allergy or allergy-induced asthma, and an  
CC infectious disease. C1C is also useful for increasing IFN-gamma, or IFN-  
CC alpha, in an individual, where the individual has idiopathic pulmonary  
CC fibrosis, or a viral infection. C1C's are useful for ameliorating a  
CC symptom of an infectious disease, or an immunoglobulin E (IgE)-related  
CC disorder in an individual, where the IgE-related disorder is allergy, or  
CC an allergy-related disorder. C1C's are also useful for treating cancer  
CC and can be used for stimulating cellular immune system cells production  
CC in an individual. This polynucleotide sequence represents a DNA sequence  
CC which is a nucleic acid moiety part of a chimeric immunomodulatory compound  
CC of the invention.

XX  
SQ Sequence 19 BP; 4 A; 2 C; 6 G; 5 T; 0 U; 2 Other;

Query Match 81.0%; Score 17; DB 8; Length 19;  
Best Local Similarity 89.5%; Pred. No. 34;  
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGACGTTGAGATG 19  
| | | | | | | | | | | | | | | | | | |  
Db 1 TNGTNGACGTTGAGATG 19

## RESULT 15

ABQ75165  
ID ABQ75165 standard; DNA; 18 BP.

XX  
AC ABQ75165;

DT 05-NOV-2002 (first entry)

XX  
DE ISS immunomodulatory oligonucleotide SEQ ID NO.14.

XX  
KM Immunostimulatory sequence; ISS; immunomodulatory; immune response;  
KM allergy; asthma; infectious disease; interferon-gamma; IFN-gamma;  
KM idiopathic pulmonary fibrosis; viral infection; mycobacterial disease;  
KM malaria; leishmaniasis; toxoplasmosis; schistosomiasis; clonorchiasis;  
KM immunoglobulin E; IgE-related disorder; anti-allergic; antiaesthetic;  
KM virucide; antibacterial; protozoacide; ss.

XX  
OS Synthetic.

XX  
PN WO200252002-A2.

XX  
PD 04-JUL-2002.

XX  
PF 27-DEC-2001; 2001WO-US050821.

XX  
PR 27-DEC-2000; 2000US-0258675P.

XX  
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX  
PI Fearon KL, Dina D;

XX  
DR WPI; 2002-657426/70.

XX  
PT Immunomodulatory polynucleotide for modulating an immune response in a  
PT subject suffering from disorders associated with Th2-type immune  
PT response, e.g. allergy, or infectious disease, comprises an  
PT immunostimulatory sequence.  
XX  
PS Example 1; Page 20; 95pp; English.

CC The present invention describes an immunomodulatory polynucleotide (1)  
CC comprising an immunostimulatory sequence (ISS). Also described: (1) an  
CC immunomodulatory composition comprising (1); (2) an immunomodulatory  
CC polynucleotide/microcarrier (IMP/MC) complex, comprising (1) linked to a  
CC biodegradable MC, where the MC is less than 10 micrometre in size; and

(3) a kit comprising (1). (1) has anti-allergic, antiasthmatic, virucide, antibacterial and protozoacide activities, and can be used as a modulator of immune response. (1) is useful for modulating an immune response in an individual suffering from disorders associated with a Th2-type immune response, especially an allergy or asthma, or an infectious disease. (1) is also useful for increasing interferon-gamma (IFN-gamma) in an individual having idiopathic pulmonary fibrosis, or IFN-alpha in an individual having a viral infection. (1) is further useful for ameliorating a symptom of an infectious disease caused by a cellular pathogen such as mycobacterial disease, malaria, leishmaniasis, toxoplasmosis, schistosomiasis and clonorchiasis in an individual, or a symptom of an immunoglobulin E (IGE)-related disorder, preferably an allergy-related disorder, in particular asthma in an individual. The present sequence represents an immunomodulatory oligonucleotide from the present invention.

Sequence 18 BP; 4 A; 4 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 6; Length 18;  
Best Local Similarity 94.4%; Pred. No. 71;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

2 CGTCGAACGTTGAGATG 19  
1 CTTGCAACGTTGAGATG 18

RESULT 16  
ADB88833  
ID ADB88833 standard; DNA; 18 BP.  
AC ADB88833;  
XX  
DT 04-DEC-2003 (first entry)  
XX  
DE Chimeric immunomodulatory compound DNA sequence, SEQ ID No 36.  
XX  
KW chimeric immunomodulatory compound; C1C; immunomodulatory activity;  
KW spacer moiety; linear hexaethylene glycol structure; HEG; immune;  
KW Th2-type; allergy; allergy-induced asthma; infectious disease; IFN-gamma;  
KW IFN-alpha; idiopathic pulmonary fibrosis; viral infection; ameliorating;  
KW immunoglobulin E; IGE; allergy; cancer;  
KW stimulating cellular immune system cell; ss.  
XX  
OS Synthetic.  
XX  
PN WO200300922-A2.  
XX  
PD 03-JAN-2003.  
XX  
PF 21-JUN-2002; 2002WO-US020025.  
XX  
PR 21-JUN-2001; 2001US-0299883P.  
PR 23-APR-2002; 2002US-0375253P.  
XX  
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
XX  
PI Fearon KL, Dina D, Tuck SF;  
XX  
DR WPI; 2003-210159/20.  
XX  
PT Novel chimeric immunomodulatory compound having immunomodulatory  
PT activity, useful for modulating an immune response and for treating  
PT cancer, has nucleic acid moieties and non-nucleic acid spacer moieties.  
XX  
PS Disclosure; Page 33; 224pp; English.  
XX  
CC The invention relates to a novel chimeric immunomodulatory compound (C1C)  
CC having immunomodulatory activity, comprising two or more nucleic acid  
CC moieties and one or more non-nucleic acid spacer moieties, where at least  
CC one non-nucleic acid spacer moiety is covalently joined to two nucleic  
CC acid moieties, where the spacer is not a polypeptide, and at least one  
CC nucleic acid moiety comprises the sequence 5'-CG-3'. The chimeric

immunomodulatory compounds more specifically contain the nucleic acid spacer moieties of linear hexaethylene glycol structure (HEG) subunits. C1C's are useful for modulating an immune response in an individual, where the individual suffers from a disorder associated with a Th2-type immune response which is an allergy or allergy-induced asthma, and an infectious disease. C1C is also useful for increasing IFN-gamma, or IFN-alpha, in an individual, where the individual has idiopathic pulmonary fibrosis, or a viral infection. C1C's are useful for ameliorating a symptom of an infectious disease, or an immunoglobulin E (IGE)-related disorder in an individual, where the IGE-related disorder is allergy, or an allergy-related disorder. C1C's are also useful for treating cancer and can be used for stimulating cellular immune system cells production in an individual. This polynucleotide sequence represents a DNA sequence which is a nucleic acid moiety part of a chimeric immunomodulatory compound of the invention.

Sequence 18 BP; 4 A; 4 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 8; Length 18;  
Best Local Similarity 94.4%; Pred. No. 71;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

2 CGTCGAACGTTGAGATG 19  
1 CTTGCAACGTTGAGATG 18

RESULT 17  
AAV80098/C  
ID AAV80098 standard; DNA; 23 BP.  
XX  
AC AAV80098;  
XX  
DT 12-MAR-1999 (first entry)  
XX  
DE Immunomodulatory oligo comprising an ISS sequence.  
XX  
KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;  
KW ISS; cancer; allergy; asthma; hepatitis B infection; papillomavirus;  
KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; ss;  
KW B. pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.  
XX  
OS Synthetic.  
XX  
PN WO9855495-A2.  
XX  
PD 10-DEC-1998.  
XX  
PF 05-JUN-1998; 98WO-US011578.  
XX  
PR 06-JUN-1997; 97US-0048793P.  
XX  
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
XX  
PI Schwartz D, Roman M, Dina D;  
XX  
DR WPI; 1999-059898/05.  
XX  
PT Immunostimulatory oligonucleotides regulate the immune system - and  
PT contain an immune-stimulating octanucleotide sequence; for treating  
PT cancer, allergic and infectious diseases.  
XX  
PS Claim 6; Page 29; 63pp; English.  
XX  
CC The invention relates to immunomodulatory oligonucleotides that comprise  
CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS  
CC sequences are selected from the group consisting of AACGTTCC, AACGTTCG,  
CC GACGTTCC, and GACGTTCC. The immunomodulatory sequences are used to treat  
CC patients needing immune regulation, such as those suffering from cancer,  
CC an allergic disease and asthma. They are also used to prevent infectious  
CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency  
CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and  
CC Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and

CC Schistosoma. The immunomodulatory sequences are used to screen for human  
CC immunostimulatory activity by incubating macrophage cells and the  
CC oligonucleotide; and determining the relative amount of Th1-biased  
CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent  
CC specific claimed examples of such immunomodulatory oligonucleotides  
XX  
SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 2; Length 23;  
Best Local Similarity 94.4%; Pred. No. 72;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGAGATGA 20  
DB 18 GTGGAACGTTGAGATGA 1

RESULT 18  
AAA38067/C  
ID AAA38067 standard; DNA; 23 BP.

AC AAA38067;  
DT 24-AUG-2000 (first entry)

DE Immunostimulatory sequence (ISS) #3.

XX Immunostimulatory sequence; ISS; immunomodulator; glycoprotein 120;  
KW gp120; human immunodeficiency virus; HIV; immune response; infection;  
KW development; ss.

OS Synthetic.

XX WO200021556-A1.

XX 20-APR-2000.

XX 08-OCT-1999; 99WO-US023677.

XX 09-OCT-1998; 98US-0103733P.

XX 07-OCT-1999; 99US-00415186.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX Tigue H, Raz E, Schwartz D, Takabayashi K;

XX WPI; 2000-317846/27.

XX Anti-HIV composition comprises immunostimulatory polynucleotides and HIV  
PT glycoprotein gp120 useful for modulating, stimulating an immune response  
PT against HIV in an HIV infected individual.

XX Disclosure; Page 16; 65pp; English.

CC The present invention relates to an immunostimulatory composition  
CC comprising a human immunodeficiency virus (HIV) antigen, and an  
CC immunomodulatory polynucleotide comprising an immunostimulatory sequence  
CC (ISS). This sequence represents an ISS that can be used in the  
CC composition. An immunostimulatory composition which comprises a gp120  
CC conjugated to an immunomodulatory polynucleotide, or is proximately  
CC associated to it and not conjugated, is used for modulating or  
CC stimulating a specific immune response against gp120 in an individual by  
CC producing anti-gp120 antibodies or gp120 specific cytotoxic T cells. It  
CC is also used for suppressing or delaying development of HIV infection in  
CC an individual infected with HIV or an individual at risk of infection  
CC with HIV, respectively. It is also used for treating an individual  
CC infected with HIV in need of immune modulation

XX Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 3; Length 23;  
Best Local Similarity 94.4%; Pred. No. 72;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGAGATGA 20  
DB 18 GTGGAACGTTGAGATGA 1

RESULT 19  
AAH75994/C  
ID AAH75994 standard; DNA; 23 BP.

AC AAH75994;

DT 15-NOV-2001 (first entry)

DE Immunomodulatory oligonucleotide #3.

XX Immunomodulatory; immunostimulatory; Th1-type immune response;  
KW Th2-type immune response; interferon; idiopathic pulmonary fibrosis;  
KW viral infection; ss.

OS Synthetic.

XX WO200168143-A2.

XX 20-SEP-2001.

XX 12-MAR-2001; 2001WO-US007843.

XX 10-MAR-2000; 2000US-0188557P.

XX 09-MAR-2001; 2001US-00802376.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX Van Nest G, Tuck S;

XX WPI; 2001-582389/65.

XX Immunomodulatory polynucleotide/microcarrier complexes comprise an  
PT immunostimulatory sequence containing polynucleotide linked to a  
PT nonbiodegradable microcarrier.

XX Disclosure; Page 18; 61pp; English.

CC The present invention relates to immunomodulatory polynucleotide/  
CC microcarrier complexes. The complexes comprise an immunostimulatory  
CC sequence (ISS), e.g. the present sequence, linked to a nonbiodegradable  
CC microcarrier provided that if the microcarrier is gold, latex or magnetic  
CC then the linkage is not biotin/avidin. The complex is useful for  
CC modulating an immune response (especially stimulating a Th1-type response  
CC or suppressing a Th2-type response), increasing interferon-gamma  
CC (especially in a patient suffering from idiopathic pulmonary fibrosis),  
CC increasing interferon-alpha (especially in patients suffering from viral  
CC infection) and reducing levels of IgE

XX Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 4; Length 23;  
Best Local Similarity 94.4%; Pred. No. 72;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGAGATGA 20  
DB 18 GTGGAACGTTGAGATGA 1

RESULT 20  
AAF77042/C  
ID AAF77042 standard; DNA; 23 BP.

AC AAF77042;

DT 15-MAY-2001 (first entry)

DE Immunostimulatory DNA #2.  
 XX Modulate; immune; antigen; immunostimulatory; ds.  
 XX Synthetic.  
 XX WO200112223-A2.  
 XX PD 22-FEB-2001.  
 XX PF 18-AUG-2000; 2000MO-US022835.  
 XX PR 19-AUG-1999; 99US-0149768P.  
 XX PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
 XX PI Van Nest G;  
 XX DR WPI; 2001-211136/21.  
 XX PT Modulating immune response to a second antigen in humans involves  
 PT administering an immunostimulatory polynucleotide comprising an  
 PT immunostimulatory sequence and a first antigen.  
 XX PS Disclosure; Page 15; 63pp; English.  
 XX CC The present invention relates to modulating an immune response to a  
 CC second antigen in an individual, involving administering to the  
 CC individual an immunomodulatory polynucleotide comprising an  
 CC immunostimulatory sequence (ISS) and a first antigen  
 XX SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;  
 QY Query Match 78.1%; Score 16.4; DB 4; Length 23;  
 Best Local Similarity 94.4%; Pred. No. 72;  
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 DB 3 GTGGAACGTTCCGAGATGA 20  
 18 GTGGAACGTTCCGAGATGA 1  
 RESULT 21  
 AAH41575/C  
 ID AAH41575 standard; DNA; 23 BP.  
 XX AC AAH41575;  
 XX DT 24-AUG-2001 (first entry)  
 XX DE Immunostimulatory sequence (ISS) SEQ ID NO:3.  
 XX OS Synthetic.  
 XX PN WO200135991-A2.  
 XX PD 25-MAY-2001.  
 XX PF 15-NOV-2000; 2000MO-US031385.  
 XX PR 15-NOV-1999; 99US-0165467P.  
 XX PR 14-NOV-2000; 2000US-00713136.  
 XX PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
 XX PI Tuck S, Van Nest G;  
 XX DR WPI; 2001-329209/34.

PT Populations of conjugate molecules comprising polynucleotide  
 PT immunostimulatory sequences polynucleotides and antigens, useful for  
 PT controlling immune responses.  
 XX PS Disclosure; Page 30; 97pp; English.  
 XX CC The present invention describes immunomodulatory populations (I) and  
 CC (II) of conjugate molecules (Cms) comprising immunostimulatory sequences  
 CC (ISS) of polynucleotides and antigens. The extent of conjugation affects  
 CC the immunological properties (e.g. the extent of antigen-specific  
 CC antibody formation, including Th1-associated antibody formation) so the  
 CC conjugates are used for altering the type and extent of immune response.  
 CC (I) and (II) have immunomodulatory, immunosuppressive and anti-allergic  
 CC activities, and can be used in the modulation of immune responses via the  
 CC stimulation of Th1 lymphocytes and Th1-associated cytokines, and  
 CC suppression of Th2 lymphocytes and cytokines. The populations (I) and  
 CC (II) of conjugate molecules may be used for modulating immune responses  
 CC in individuals e.g. for the treatment of an allergic condition. (I) and  
 CC (II) may be used to modulate immune responses and therefore prevent  
 CC potentially harmful reactions to antigens. The present sequence  
 CC represents an ISS polynucleotide which is used in the exemplification of  
 XX the present invention  
 SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;  
 QY Query Match 78.1%; Score 16.4; DB 5; Length 23;  
 Best Local Similarity 94.4%; Pred. No. 72;  
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 DB 3 GTGGAACGTTCCGAGATGA 20  
 18 GTGGAACGTTCCGAGATGA 1  
 RESULT 22  
 AAS14666/C  
 ID AAS14666 standard; DNA; 23 BP.  
 XX AC AAS14666;  
 XX DT 18-DEC-2001 (first entry)  
 XX DE Immunostimulatory sequence, ISS #3.  
 XX OS Synthetic.  
 XX PN WO200168116-A2.  
 XX PD 20-SEP-2001.  
 XX PF 12-MAR-2001; 2001MO-US007839.  
 XX PR 10-MAR-2000; 2000US-0188583P.  
 XX PR 09-MAR-2001; 2001US-00802686.  
 XX PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
 XX PI Van Nest G;  
 XX DR WPI; 2001-607438/69.  
 XX PT Suppressing a respiratory syncytial virus infection by administering an  
 PT immunostimulatory sequence at the site of infection is useful to prevent  
 PT and treat lower respiratory tract viral infections.  
 XX PS Disclosure; Page 15; 40pp; English.

CC The invention relates to suppressing a respiratory syncytial virus (RSV)  
CC infection in an exposed individual, comprising administering a  
CC polynucleotide comprising an immunostimulatory sequence (ISS) comprising  
CC the sequence 5'-C, G-3', where an RSV antigen is not administered. The  
CC invention is used to prevent and treat respiratory syncytial virus  
CC infection of the lower respiratory tract and other viruses including  
CC influenza virus, rhinovirus, adenovirus, measles virus, mumps virus,  
CC parainfluenza virus, rubella virus, poxvirus, parvovirus, hantavirus and  
CC varicella virus. A kit for carrying out the administration is also  
CC included unlike the prior art antiviral agent ribavirin, which is a  
CC potential teratogen, the invention provides a treatment which does not  
CC carry unacceptable side effects. Other prior art medicaments treat the  
CC symptoms only, whilst the invention treats the infection. The present  
CC sequence is an ISS of the invention

XX  
SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 5; Length 23;  
Best Local Similarity 94.4%; Pred. No. 72;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 GTCGACGTCGAGATGA 20  
Db 18 GTGGAACGTCGAGATGA 1

RESULT 23  
ABA03835/C  
ID ABA03835 standard; DNA; 23 BP.

AC ABA03835;  
XX  
DT 12-FEB-2002 (first entry)  
XX  
DE Immunostimulatory sequence (ISS) SEQ ID NO:3.

XX  
XX Immunomodulatory polynucleotide/microcarrier complex; IMP/MC; ISB;  
XX immunomodulation; immunostimulation; phosphorothioate; immunomodulator;  
XX antiallergic; antibacterial; antiparasitic; antiparasitic; hepatotropic;  
XX nephrotoxic; interferon-alpha stimulator; interferon-gamma stimulator;  
XX immunoglobulin E stimulator; immune response; IPF; scleroderma; malaria;  
XX idiopathic pulmonary fibrosis; cutaneous radiation-induced fibrosis;  
XX hepatic fibrosis; renal fibrosis; infectious disease; leishmaniasis;  
XX mycobacterial disease; toxoplasmosis; schistosomiasis; chionorchiasis;  
XX allergy; allergy-induced asthma; prophylactic vaccine; cancer; ss.

OS Synthetic.  
XX  
XX WO200168144-A2.  
XX  
XX 20-SEP-2001.  
XX  
XX 12-MAR-2001; 2001WO-US007848.  
XX  
XX 10-MAR-2000; 2000US-0188303P.  
XX  
XX 09-MAR-2001; 2001US-00802359.  
XX  
XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
XX  
XX Van Nest G, Tuck S;  
XX  
XX WPI; 2002-049002/06.  
XX  
XX New immunomodulatory polynucleotide/microcarrier complex, useful for  
XX modulating the immune response of individuals, particularly humans, or  
XX for treating idiopathic pulmonary fibrosis, scleroderma, malaria or  
XX allergies.  
XX  
XX Disclosure; Page 18; 63pp; English.  
XX  
XX The present invention describes an immunomodulatory polynucleotide/  
XX microcarrier (IMP/MC) complex (I), which comprises a polynucleotide  
XX having an immunostimulatory sequence (ISS) linked to a biodegradable

CC microcarrier (MC). The ISS comprises the sequence: 5'-CG-3', where the MC  
CC is less than 10 micro m in size. (I) has immunomodulator, antiallergic,  
CC antibacterial, antiparasitic, antiparasitic, hepatotropic and  
CC nephrotoxic activities. It can be used as an interferon (IFN)-alpha  
CC stimulator, IFN-gamma stimulator or an immunoglobulin E (ISB) stimulator.  
CC (I) can be used for modulating the immune response of individuals,  
CC particularly humans. The IMP/MC complex is particularly useful for  
CC treating idiopathic pulmonary fibrosis (IPF), scleroderma, cutaneous  
CC radiation-induced fibrosis, hepatic fibrosis, infectious diseases caused by  
CC induced hepatic fibrosis, renal fibrosis, infectious diseases caused by  
CC cellular pathogen (e.g. a mycobacterial disease, malaria, leishmaniasis,  
CC toxoplasmosis, schistosomiasis or chionorchiasis), or disorders  
CC associated with a Th2-type immune response (e.g. allergies or allergy-  
CC induced asthma). The IMP/MC may also be used in individuals receiving  
CC therapeutic or prophylactic vaccines, in individuals suffering from  
CC cancer, or in individuals at risk of exposure to an infectious agent. The  
CC present sequence represents an ISS given in the exemplification of the  
CC present invention

XX  
SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 72;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 GTCGACGTCGAGATGA 20  
Db 18 GTGGAACGTCGAGATGA 1

RESULT 24  
ABA03846/C  
ID ABA03846 standard; DNA; 23 BP.

AC ABA03846;  
XX  
DT 12-FEB-2002 (first entry)  
XX  
DE Immunostimulatory sequence (ISS) SEQ ID NO:3.

XX  
XX Immunostimulatory sequence; ISS; immunostimulation; viral infection;  
XX immunomodulation; virucide; gene therapy; viraemia; ss.

OS Synthetic.  
XX  
XX WO200168077-A2.  
XX  
XX 20-SEP-2001.  
XX  
XX 12-MAR-2001; 2001WO-US007840.  
XX  
XX 10-MAR-2000; 2000US-0188302P.  
XX  
XX 09-MAR-2001; 2001US-00802685.  
XX  
XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
XX  
XX Van Nest G;  
XX  
XX WPI; 2002-048999/06.  
XX  
XX Reducing severity, recurrence or duration of symptom of virus infection,  
XX or reducing viraemia or blood levels of virus antigen, comprises  
XX administering a polynucleotide having an immunostimulatory sequence.  
XX  
XX Disclosure; Page 20; 65pp; English.  
XX  
XX The present invention describes a method for reducing severity of a  
XX symptom of virus infection in an individual infected with a virus. The  
XX method comprises administering a composition consisting of a  
XX polynucleotide having an immunostimulatory sequence (ISS). The ISS  
XX comprises the sequence 5'-C, G, pyrimidine, pyrimidine, C, G-3'. An antigen is  
XX administered in conjunction with the composition. ISS has virucide  
XX activity and can be used in gene therapy. The method using the ISS can be

CC used for suppressing, ameliorating and/or preventing viral infections to  
CC an individual who may be at risk of being exposed to, exposed to or  
CC infected by a virus. It may also be used in reducing the recurrence or  
CC duration of a symptom of viral infection, delaying the development of a  
CC virus infection, and reducing viraemia or blood levels of virus antigens.  
CC The present sequence represents an ISS given in the exemplification of  
CC the present invention

XX  
SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 72;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGCAGATGA 20  
|||  
18 GTGGAACGTTGCAGATGA 1

DB 18 GTGGAACGTTGCAGATGA 1

#### RESULT 25

AA516339/C  
ID AA516339 standard; DNA; 23 BP.

XX  
AC AA516339;

XX  
DT 14-FEB-2002 (first entry)

DE ISS polynucleotide #3 useful for treating herpes virus infections.

XX Herpes simplex virus; HSV infection; immunostimulatory sequence; ISS;

KW immune response; alphaherpesvirinae; herpes virus zoster virus; VZV;

KM HSV-1; HSV-2; chicken pox; herpes labialis; cold sore; genital herpes;

KM virucide; ss.

XX  
OS Synthetic.

XX  
PN WO200168103-A2.

XX  
PD 20-SEP-2001.

XX  
PF 12-MAR-2001; 2001WO-US007841.

XX  
PR 10-MAR-2000; 2000US-0188556P.

XX  
PR 09-MAR-2001; 2001US-00802518.

XX  
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX  
PI Van Nest G;

XX  
DR WPI; 2002-041171/05.

XX  
PT Preventing, reducing the severity or reducing the recurrence of an

PT infection or symptom of herpes simplex virus (HSV), e.g. HSV-2, comprises

PT administering an immunostimulatory sequence to an individual.

XX  
PS Disclosure; Page 19; 49pp; English.

XX  
CC The present invention relates to novel methods of treating, preventing,

CC or reducing the severity or recurrence of a symptom of herpes simplex

CC virus (HSV) infection in an individual who has been exposed to or who is

CC infected with HSV. The method comprises administering a polynucleotide

CC having an immunostimulatory sequence (ISS; AA516337-AA516345) which

CC induces an immune response. A composition containing ISS is administered

CC without a HSV (alphaherpesvirinae) antigen. The composition can be

CC included in a kit for ameliorating or preventing a symptom of HSV

CC infection caused by herpes virus zoster virus (VZV), HSV-1 and

CC labialis (cold sores) and genital herpes. The present sequence represents

CC one of the ISS polynucleotides of the invention. Note: The present

CC sequence is shown as single stranded in the specification, but the

CC patentees state on page 20 that this sequence may be double stranded

XX  
SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 6; Length 23;  
Best Local Similarity 94.4%; Pred. No. 72;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGCAGATGA 20  
|||  
18 GTGGAACGTTGCAGATGA 1

DB 18 GTGGAACGTTGCAGATGA 1

#### RESULT 26

AA516350/C  
ID AA516350 standard; DNA; 23 BP.

XX  
AC AA516350;

XX  
DT 14-FEB-2002 (first entry)

DE ISS polynucleotide #3 useful for treating papillomavirus infections.

XX Animal papillomavirus infection; human papillomavirus; HPV; STD; wart;

KW sexually transmitted disease; cervical cancer; immune response;

KM immunostimulatory sequence; ISS; virucide; ss.

XX  
OS Synthetic.

XX  
PN WO200168117-A2.

XX  
PD 20-SEP-2001.

XX  
PF 12-MAR-2001; 2001WO-US007842.

XX  
PR 10-MAR-2000; 2000US-0188265P.

XX  
PR 09-MAR-2001; 2001US-00802445.

XX  
PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX  
PI Van Nest G;

XX  
DR WPI; 2002-041172/05.

XX  
PT Treating, preventing or ameliorating papillomavirus infections, comprises

PT administering a composition comprising a polynucleotide having an

PT immunostimulatory sequence to the individual.

XX  
PS Disclosure; Page 20; 44pp; English.

XX  
CC The present invention relates to novel methods of treating, preventing,

CC or reducing the severity or recurrence of a symptom of papillomavirus

CC infection in an individual that has been exposed to or who is infected

CC with papillomavirus. The method comprises administering a polynucleotide

CC having an immunostimulatory sequence (ISS; AA516348-AA516355) which

CC induces an immune response. A composition containing ISS is administered

CC without a papillomavirus antigen. The composition can be included in a

CC kit for ameliorating or preventing a symptom of human or animal

CC papillomavirus infection. Infections with human papillomavirus (HPV)

CC which can be prevented or treated using the method of the invention

CC include sexually transmitted diseases (STDs), warts, papillomas and

CC cervical cancer. The present sequence represents one of the ISS

CC polynucleotides of the invention. Note: The present sequence is shown as

CC single stranded in the specification, but the patentees state on page 20

CC that this sequence may be double stranded

XX  
SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 6; Length 23;

Best Local Similarity 94.4%; Pred. No. 72;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 GTCGAACGTTGCAGATGA 20  
|||  
18 GTGGAACGTTGCAGATGA 1

DB 18 GTGGAACGTTGCAGATGA 1

## RESULT 27

ABR03858/c  
ID ABA03858 standard; DNA; 23 BP.

XX ABA03858;

XX 12-FEB-2002 (first entry)

XX Immunostimulatory sequence (ISS) SEQ ID NO:3.

XX Immunostimulatory sequence; ISS; immunomodulation; HBV; HCV; infection;

XX hepatitis B virus; hepatitis C virus; virucide; anti-inflammatory;

XX hepatotropic; gene therapy; hepatitis infection; viraemia; jaundice;

XX fatigue; abdominal pain; portal hypertension; cirrhosis; ss.

XX Synthetic.

XX WO200168078-A2.

XX 20-SEP-2001.

XX 12-MAR-2001; 2001WO-US007931.

XX 10-MAR-2000; 2000US-0188301P.

XX 09-MAR-2001; 2001US-00802370.

XX (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX Van Nest G;

XX WPI; 2002-049000/06.

XX Reducing viraemia and blood levels of hepatitis virus antigen in an

XX individual infected with hepatitis B virus, comprises administering a

XX composition comprising a polynucleotide having an immunostimulatory

XX sequence.

XX Disclosure; Page 20; 43pp; English.

XX The present invention describes a method for reducing viraemia or blood

XX levels of a hepatitis virus antigen in an individual infected with

XX hepatitis B virus (HBV). The method comprises administering a composition

XX comprising a polynucleotide having an immunostimulatory sequence (ISS) to

XX the individual, where the ISS comprises the sequence 5'-C<sub>1</sub>-G<sub>1</sub>3', an HBV

XX antigen is not administered in conjunction with administration of the

XX composition, and where the composition is administered in an amount

XX sufficient to reduce HBV viraemia or blood levels of a hepatitis virus

XX antigen. ISS has virucide, anti-inflammatory and hepatotropic activities,

XX and/or ameliorating hepatitis infection in an individual, especially for

XX preventing, palliating, ameliorating, reducing and/or eliminating one or

XX more symptoms of HBV or HCV (hepatitis C virus) infection without

XX administering HBV or HCV antigens. The method is specifically useful for

XX reducing viraemia and hepatitis viral antigen in blood. ISS-containing

XX polynucleotides may also be used to improve physical symptoms such as

XX jaundice, fatigue, abdominal pain, and other clinical/laboratory

XX findings associated with hepatitis such as blood levels of liver enzymes,

XX portal hypertension, or cirrhosis. The present sequence represents an ISS

XX oligonucleotide given in the exemplification of the present invention

SQ Sequence 23 BP; 6 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

Query Match 78.1%; Score 16.4; DB 6; Length 23;

Best Local Similarity 94.4%; Pred. No. 72;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GTGACGCTTCGAGATGA 20

Db 18 GTGACGCTTCGAGATGA 1

RESULT 28

ABL12595/c  
ID ABL12595 standard; cDNA; 1017 BP.

XX ABL12595;

XX 26-MAR-2002 (first entry)

XX Drosophila melanogaster expressed polynucleotide SEQ ID NO 32267.

XX Drosophila; developmental biology; cell signalling; insecticide;

XX pharmaceutical; gene; ss.

XX Drosophila melanogaster.

XX WO200171042-A2.

XX 27-SEP-2001.

XX 23-MAR-2001; 2001WO-US009231.

XX 23-MAR-2000; 2000US-0191637P.

XX 11-JUL-2000; 2000US-00614150.

XX (PEKE) PE CORP NY.

XX Venter JC, Adams M, Li PWD, Myers EW;

XX WPI; 2001-65860/75.

XX P-PSDB; ABB68492.

XX New isolated nucleic acid detection reagent for detecting 1000 or more

XX genes from Drosophila and for elucidating cell signaling and cell-cell

XX interactions.

XX Claim 1; SEQ ID NO 32267; 21pp + Sequence Listing; English.

XX The invention relates to an isolated nucleic acid detection reagent

XX capable of detecting 1000 or more genes from Drosophila. The invention is

XX useful in developmental biology and in elucidating cell signaling and

XX cell-cell interactions in higher eukaryotes for the development of

XX insecticides, therapeutics and pharmaceutical drugs. The invention

XX discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA

XX sequences (ABL10840-ABL16175) and the encoded proteins (AB857737-

XX AB872072). The sequence data for this patent did not form part of the

XX printed specification, but was obtained in electronic format directly

XX from WIPO at ftp.wipo.int/pub/published\_pct\_sequences

SQ Sequence 1017 BP; 246 A; 275 C; 230 G; 206 T; 0 U; 0 Other;

Query Match 77.1%; Score 16.2; DB 4; Length 1017;

Best Local Similarity 85.7%; Pred. No. 1.2e+02;

Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TCGTCGACGCTTCGAGATGAT 21

Db 253 TCGTCGACGCTTCGAGATGAT 233

## RESULT 29

ACA47671  
ID ACA47671 standard; DNA; 1020 BP.

XX ACA47671;

XX 19-JUN-2003 (first entry)

XX Prokaryotic essential gene #29328.

XX Antisense; ds; prokaryotic essential gene; cell proliferation;

XX drug design; gene.

XX Staphylococcus haemolyticus.

XX



PN WO20027183-A2.  
 XX 03-OCT-2002.  
 XX 21-MAR-2002; 2002WO-US009107.  
 XX 21-MAR-2001; 2001US-00815242.  
 PR 06-SEP-2001; 2001US-00948993.  
 PR 25-OCT-2001; 2001US-00342923P.  
 PR 08-FEB-2002; 2002US-00072851.  
 PR 06-MAR-2002; 2002US-0362699P.  
 XX (ELIT-) ELITRA PHARM INC.  
 XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW,  
 PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;  
 XX WPI; 2003-029926/02.  
 DR P-PSDB; ABU43801.  
 XX New antisense nucleic acid, useful for identifying proteins or screening  
 PT for homologous nucleic acids required for cellular proliferation to  
 PT isolate candidate molecules for rational drug discovery programs.  
 XX Claim 14; SEQ ID NO 35541; 1766pp; English.  
 XX The invention relates to an isolated nucleic acid comprising any one of  
 CC the 6213 antisense sequences given in the specification where expression  
 CC of the nucleic acid inhibits proliferation of a cell. Also included are:  
 CC (1) a vector comprising a promoter operably linked to the nucleic acid  
 CC encoding a polypeptide whose expression is inhibited by the antisense  
 CC nucleic acid; (2) a host cell containing the vector; (3) an isolated  
 CC polypeptide or its fragment whose expression is inhibited by the  
 CC antisense nucleic acid; (4) an antibody capable of specifically binding  
 CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular  
 CC proliferation or the activity of a gene in an operon required for  
 CC proliferation; (7) identifying a compound that influences the activity of  
 CC the gene product or that has an activity against a biological pathway  
 CC required for proliferation, or that inhibits cellular proliferation; (8)  
 CC identifying a gene required for cellular proliferation or the biological  
 CC pathway in which a proliferation-required gene or its gene product lies  
 CC or a gene on which the test compound that inhibits proliferation of an  
 CC organism acts; (9) manufacturing an antibiotic; (10) profiling a  
 CC compound's activity; (11) a culture comprising strains in which the gene  
 CC product is overexpressed or underexpressed; (12) determining the extent  
 CC to which each of the strains is present in a culture or collection of  
 CC strains; or (13) identifying the target of a compound that inhibits the  
 CC proliferation of an organism. The antisense nucleic acids are useful for  
 CC identifying proteins or screening for homologous nucleic acids required  
 CC for cellular proliferation to isolate candidate molecules for rational  
 CC drug discovery programs, or for screening homologous nucleic acids  
 CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,  
 CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target  
 CC prokaryotic essential genes. Note: The sequence data for this patent did  
 CC not form part of the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences  
 XX  
 SQ Sequence 1020 BP; 351 A; 148 C; 204 G; 317 T; 0 U; 0 Other;  
 Query Match 77.1%; Score 16.2; DB 7; Length 1020;  
 Best Local Similarity 85.7%; Pred. No. 1.2e+02;  
 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

AC ABQ14749;  
 XX 12-JUL-2002 (first entry)  
 DT  
 XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 1340.  
 DE  
 XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;  
 KM drug; side effect; cancer; central nervous system; cardiovascular;  
 KM gastrointestinal; respiratory system; single nucleotide polymorphism;  
 KM SNP; cell differentiation; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 PM WO200218632-A2.  
 XX 07-MAR-2002.  
 XX  
 PF 01-SEP-2001; 2001WO-EP010074.  
 XX  
 PR 01-SEP-2000; 2000DE-01043826.  
 PR 05-SEP-2000; 2000DE-01044543.  
 XX  
 PA (EPIC-) EPIGENOMICS AG.  
 XX  
 PI Olek A, Piepenbrock C, Berlin K, Guetig D;  
 XX WPI; 2002-371829/40.  
 DR  
 XX  
 XX Determining the degree of cytosine methylation in genomic DNA, useful for  
 PT diagnosis and prognosis, comprises selective hybridization of amplicons  
 PT from chemically treated DNA.  
 XX  
 PS Claim 12; 56pp + Sequence Listing; 56pp; German.  
 XX  
 XX This invention describes a novel method for determining the degree of  
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a  
 CC genomic sample of DNA. The sample is treated chemically to convert  
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic  
 CC DNA that contains the target C is amplified to form a labeled amplicon.  
 CC The amplicon is hybridised to two classes, each with at least one member,  
 CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the  
 CC degree of hybridisation to both classes is determined from the label on  
 CC the amplicon. From the ratio of labels hybridised to the two classes of  
 CC oligomers, the degree of methylation is calculated. The method is used:  
 CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs  
 CC and of a wide range of diseases, e.g. cancer, disorders of the central  
 CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,  
 CC particularly by detecting mutations or single nucleotide polymorphisms  
 CC (SNPs); and (ii) for differentiation of cell or tissue types and for  
 CC investigating cell differentiation. The method allows the methylation  
 CC status of many C residues to be determined simultaneously. ABQ1410-  
 CC ABQ4121 represent genomic DNA sequences used to illustrate the method  
 CC for determining the degree of cytosine methylation described in the  
 CC disclosure of the invention  
 XX  
 SQ Sequence 1676 BP; 643 A; 513 C; 171 G; 349 T; 0 U; 0 Other;  
 Query Match 77.1%; Score 16.2; DB 6; Length 1676;  
 Best Local Similarity 85.7%; Pred. No. 1.2e+02;  
 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

RESULT 30  
 ABQ14749/C  
 ID ABQ14749 standard; DNA; 1676 BP.  
 XX

RESULT 31  
 ABQ14748  
 ID ABQ14748 standard; DNA; 1676 BP.  
 XX  
 AC ABQ14748;  
 XX  
 DT 12-JUL-2002 (first entry)

XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 1339.  
DE Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;  
XX drug; side effect; cancer; central nervous system; cardiovascular;  
KM gastrointestinal; respiratory system; single nucleotide polymorphism;  
XX SNP; cell differentiation; ds.  
OS Homo sapiens.  
XX WO200218632-A2.  
XX  
XX PD 07-MAR-2002.  
XX  
XX PF 01-SEP-2001; 2001WO-EP010074.  
XX  
XX PR 01-SEP-2000; 2000DE-01043826.  
XX PR 05-SEP-2000; 2000DE-01044543.  
XX  
XX PA (EPiG-) EPIGENOMICS AG.  
XX  
XX PI Olek A., Piepenbrock C., Berlin K., Guetig D;  
XX  
XX DR WPI; 2002-371829/40.  
XX  
XX PT Determining the degree of cytosine methylation in genomic DNA, useful for  
XX diagnosis and prognosis, comprises selective hybridization of amplicons  
XX from chemically treated DNA.  
XX  
XX PS Claim 12; 56bp + Sequence Listing; 56bp; German.  
XX  
XX CC This invention describes a novel method for determining the degree of  
XX methylation of a particular cytosine in a motif 5'-CpG-3', present in a  
XX genomic sample of DNA. The sample is treated chemically to convert  
XX cytosine (C) but not methylated C, to uracil, then part of the genomic  
XX DNA that contains the target C is amplified to form a labeled amplicon.  
XX The amplicon is hybridised to two classes, each with at least one member,  
XX of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the  
XX degree of hybridisation to both classes is determined from the label on  
XX the amplicon. From the ratio of labels hybridised to the two classes of  
XX oligomers, the degree of methylation is calculated. The method is used:  
XX (i) for diagnosis and/or prognosis of side effects of therapeutic drugs  
XX and of a wide range of diseases, e.g. cancer, disorders of the central  
XX nervous, cardiovascular, gastrointestinal and respiratory systems etc.,  
XX particularly by detecting mutations or single nucleotide polymorphisms  
XX (SNP's); and (ii) for differentiation of cell or tissue types and for  
XX investigating cell differentiation. The method allows the methylation  
XX status of many C residues to be determined simultaneously. ABQ13410-  
XX ABQ54121 represent genomic DNA sequences used to illustrate the method  
XX for determining the degree of cytosine methylation described in the  
XX disclosure of the invention  
XX  
SQ Sequence 1676 BP; 349 A; 171 C; 513 G; 643 T; 0 U; 0 Other;

Query Match 77.1%; Score 16.2; DB 6; Length 1676;  
Best Local Similarity 85.7%; Pred. No. 1.2e+02;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TCGTCGACGCTCGAGATGAT 21  
| | | | | | | | | | | | | | | | | | | | | |  
Db 513 TCGTTAGACGCTCGAGATGAT 533

RESULT 32  
ABL12594  
ID ABL12594 standard; cDNA; 3017 BP.  
XX  
XX ABL12594;  
AC  
XX 26-MAR-2002 (first entry)  
DT  
XX  
XX Drosophila melanogaster expressed polynucleotide SEQ ID NO 32264.  
DE  
XX

KW Drosophila; developmental biology; cell signalling; insecticide;  
KM pharmaceutical; gene; ss.  
XX  
XX OS Drosophila melanogaster.  
XX  
XX PN WO200171042-A2.  
XX  
XX PD 27-SEP-2001.  
XX  
XX PF 23-MAR-2001; 2001WO-US009231.  
XX  
XX PR 23-MAR-2000; 2000US-0191637P.  
XX PR 11-JUL-2000; 2000US-00614150.  
XX  
XX PA (PEKE ) PE CORP NY.  
XX  
XX PI Venter JC, Adams M, Li PWD, Myers EW;  
XX  
XX DR WPI; 2001-656860/75.  
XX DR P-PSDB; ABB68491.  
XX  
XX PT New isolated nucleic acid detection reagent for detecting 1000 or more  
XX genes from Drosophila and for elucidating cell signaling and cell-cell  
XX interactions.  
XX  
XX PS Claim 1; SEQ ID NO 32264; 21pp + Sequence Listing; English.  
XX  
XX CC The invention relates to an isolated nucleic acid detection reagent  
XX capable of detecting 1000 or more genes from Drosophila. The invention is  
XX useful in developmental biology and in elucidating cell signaling and  
XX cell-cell interactions in higher eukaryotes for the development of  
XX insecticides, therapeutics and pharmaceutical drugs. The invention  
XX discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA  
XX sequences (ABL01840-ABL16175) and the encoded proteins (ABB57737-  
XX ABB12072). The sequence data for this patent did not form part of the  
XX printed specification, but was obtained in electronic format directly  
XX from WIPO at ftp.wipo.int/pub/published\_pct\_sequences  
XX  
SQ Sequence 3017 BP; 808 A; 667 C; 672 G; 870 T; 0 U; 0 Other;

Query Match 77.1%; Score 16.2; DB 4; Length 3017;  
Best Local Similarity 85.7%; Pred. No. 1.3e+02;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TCGTCGACGCTCGAGATGAT 21  
| | | | | | | | | | | | | | | | | | | | | |  
Db 1765 TCGTCGACGCTCGAGATGAT 1785

RESULT 33  
ACA33960/c  
ID ACA33960 standard; DNA; 3126 BP.  
XX  
XX ACA33960;  
AC  
XX 19-JUN-2003 (first entry)  
DT  
XX  
XX DE Prokaryotic essential gene #15617.  
XX  
XX KM Antisense; ds; prokaryotic essential gene; cell proliferation;  
XX drug design; gene.  
XX  
XX OS Enterococcus faecium.  
XX  
XX PN WO200271183-A2.  
XX  
XX PD 03-OCT-2002.  
XX  
XX PF 21-MAR-2002; 2002WO-US009107.  
XX  
XX PR 21-MAR-2001; 2001US-00815242.  
XX PR 06-SEP-2001; 2001US-00948993.  
XX PR 25-OCT-2001; 2001US-0342923P.  
XX

PR 08-FEB-2002; 2002US-00072851.  
 PR 06-MAR-2002; 2002US-0362699P.  
 XX  
 PA (ELIT-) ELITRA PHARM INC.  
 XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;  
 PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;  
 XX WPI: 2003-029926/02.  
 DR P-PSDB; ABU030090.  
 PT New antisense nucleic acids, useful for identifying proteins or screening  
 PT for homologous nucleic acids required for cellular proliferation to  
 PT isolate candidate molecules for rational drug discovery programs.  
 XX  
 PS Claim 14; SEQ ID NO 21830; 1766pp; English.  
 CC The invention relates to an isolated nucleic acid comprising any one of  
 CC the 6213 antisense sequences given in the specification where expression  
 CC of the nucleic acid inhibits proliferation of a cell. Also included are:  
 CC (1) a vector comprising a promoter operably linked to the nucleic acid  
 CC encoding a polypeptide whose expression is inhibited by the antisense  
 CC nucleic acid; (2) a host cell containing the vector; (3) an isolated  
 CC polypeptide or its fragment whose expression is inhibited by the  
 CC antisense nucleic acid; (4) an antibody capable of specifically binding  
 CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular  
 CC proliferation or the activity of a gene in an operon required for  
 CC proliferation; (7) identifying a compound that influences the activity of  
 CC the gene product or that has an activity against a biological pathway  
 CC required for proliferation, or that inhibits cellular proliferation; (8)  
 CC identifying a gene required for cellular proliferation or the biological  
 CC pathway in which a proliferation-required gene or its gene product lies  
 CC or a gene on which the test compound that inhibits proliferation of an  
 CC organism acts; (9) manufacturing an antibiotic; (10) profiling a  
 CC compound's activity; (11) a culture comprising strains in which the gene  
 CC product is overexpressed or underexpressed; (12) determining the extent  
 CC to which each of the strains is present in a culture or collection of  
 CC strains; or (13) identifying the target of a compound that inhibits the  
 CC proliferation of an organism. The antisense nucleic acids are useful for  
 CC identifying proteins or screening for homologous nucleic acids required  
 CC for cellular proliferation to isolate candidate molecules for rational  
 CC drug discovery programs, or for screening homologous nucleic acids  
 CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,  
 CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target  
 CC prokaryotic essential genes. Note: The sequence data for this patent did  
 CC not form part of the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences  
 CC  
 SQ Sequence 3126 BP; 1280 A; 536 C; 627 G; 683 T; 0 U; 0 Other;  
 QY Query Match 77.1%; Score 16.2; DB 7; Length 3126;  
 DB Best Local Similarity 85.7%; Pred. No. 1.3e+02;  
 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 1 TCGTCGAACGTTGAGATGAT 21  
 2719 TCGTTAAACGTTGAGATGAT 2699  
 RESULT 34  
 ID ADG91948 standard; DNA; 3135 BP.  
 AC ADG91948;  
 XX  
 XX  
 DT 01-JAN-2004 (first entry)  
 XX  
 DE E. faecium DNA sequence SEQ ID 1575.  
 XX  
 KW ds; gene; urinary tract infection; bacteraemia; endocarditis; wound;  
 KW abdominal-pelvic infection.  
 XX

OS Enterococcus faecium.  
 XX  
 XX US6583275-B1.  
 XX  
 PD 24-JUN-2003.  
 XX  
 XX 30-JUN-1998; 98US-00107532.  
 XX  
 XX 02-JUL-1997; 97US-0051571P.  
 PR 14-MAY-1998; 98US-0085598P.  
 XX  
 XX (GENO-) GENOME THERAPEUTICS CORP.  
 XX  
 PA Doucette-Stamm LA, Bush D;  
 PI WPI: 2003-799836/75.  
 DR P-PSDB; ADG95602.  
 XX  
 PT New isolated nucleic acid derived from Enterococcus faecium encoding an  
 PT Enterococcus faecium polypeptide useful for detection, prevention and  
 PT treatment of a pathological condition resulting from a bacterial  
 PT infection.  
 XX  
 PS Example 1; SEQ ID NO 1575; 243pp; English.  
 CC The invention relates to an isolated nucleic acid derived from  
 CC Enterococcus faecium encoding an Enterococcus faecium polypeptide having  
 CC one of 10 fully defined sequences given in the (or comprising 40  
 CC sequential nucleotides chosen from any of the nucleic acids, its  
 CC complement or sequences hybridising to it). Also included are a  
 CC recombinant vector comprising the nucleic acid operably linked to  
 CC transcription regulatory element, a cell comprising the vector and a  
 CC single-stranded probe comprising the nucleic acid. The nucleic acids are  
 CC chosen from 3654 disclosed sequences encoding 3654 disclosed proteins.  
 CC The nucleic acids is useful for diagnosing pathological conditions  
 CC resulting from *E. faecium* bacterial infection (e.g. urinary tract  
 CC infection, bacteraemia, endocarditis, wounds and abdominal-pelvic  
 CC infection) and for screening drugs such as agonists and antagonists. The  
 CC nucleic acid is useful for recombinant production of *Candida albicans* -  
 CC derived peptides or antisense polypeptides. Pharmaceutical compositions  
 CC and vaccines containing the nucleic acid are useful for preventing or  
 CC treating Enterococcus faecium infections. The present sequence represents  
 CC one if the disclosed *E. faecium* nucleic acids.  
 CC  
 SQ Sequence 3135 BP; 1285 A; 537 C; 628 G; 685 T; 0 U; 0 Other;  
 QY Query Match 77.1%; Score 16.2; DB 9; Length 3135;  
 DB Best Local Similarity 85.7%; Pred. No. 1.3e+02;  
 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 1 TCGTCGAACGTTGAGATGAT 21  
 2725 TCGTTAAACGTTGAGATGAT 2705  
 RESULT 35  
 ID AAV17086/c  
 XX AAV17086 standard; cDNA; 4024 BP.  
 AC AAV17086;  
 XX  
 XX 21-JUL-1998 (first entry)  
 DT  
 XX  
 DE Bos taurus tubulin-folding cofactor D gene.  
 XX  
 XX Tubulin folding; cofactor; alpha-tubulin; beta-tubulin; unfolded; folded;  
 KW treatment; hyper-proliferative diseases; cancer; gout; ss.  
 XX  
 OS Bos taurus.  
 XX  
 XX  
 FH Key Location/Qualifiers  
 FT 192..3791  
 FT CDS  
 FT /\*tag= a

FT /product= "bovine tubulin-folding cofactor D"  
 XX WO9804587-A1.  
 FN 05-FEB-1998.  
 PD XX  
 PF 25-JUL-1997; 97WO-US014076.  
 XX 25-JUL-1996; 96US-0023089P.  
 PR (UVMY ) UNIV NEW YORK STATE.  
 PA XX  
 PI Cowan NJ;  
 XX WPI; 1998-130618/12.  
 DR P-PSDB; AAM47206.  
 XX  
 PT New isolated cofactor(s) for tubulin folding - are useful as targets for  
 PT identifying agents which interfere with folding in the treatment of hyper-  
 PT -proliferative diseases such as cancer.  
 PS  
 PS Claim 4; Page 57-61; 87pp; English.  
 CC The sequence is that encoding bovine tubulin-folding cofactor D which may  
 CC be useful as a target for interfering with the production of productively  
 CC folded alpha- and beta-tubulins. Since tubulin function is essential for  
 CC cell division and proliferation, agents which interfere with tubulin  
 CC function can serve as useful antiproliferative compounds. Such  
 CC interfering agents have potential utility as agents for the treatment of  
 CC hyperproliferative diseases such as cancer and the treatment of gout  
 XX  
 SQ Sequence 4024 BP; 752 A; 1209 C; 1266 G; 797 T; 0 U; 0 Other;  
 Query Match 77.1%; Score 16.2; DB 2; Length 4024;  
 Best Local Similarity 85.7%; Pred. No. 1.3e+02;  
 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 QY 1 TCCTCGACGCTTCGAGATGAT 21  
 DB 557 TCCTCGACGCTTCGAGATGAT 537  
 RESULT 36  
 ID ADB88936 standard; DNA; 66 BP.  
 AC ADB88936;  
 XX  
 DT 04-DEC-2003 (first entry)  
 XX  
 DE Chimeric immunomodulatory compound DNA sequence, SEQ ID NO 139.  
 XX  
 XX chimeric immunomodulatory compound; CIC; immunomodulatory activity;  
 KW spacer moiety; linear hexaethylene glycol structure; HEG; immune;  
 KW Th2-type; allergy; allergy-induced asthma; infectious disease; IFN-gamma;  
 KW IFN-alpha; idiopathic pulmonary fibrosis; viral infection; ameliorating;  
 KW immunoglobulin E; IgE; allergy; cancer;  
 KW stimulating cellular immune system cell; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO2003000922-A2.  
 PD 03-JAN-2003.  
 XX  
 PF 21-JUN-2002; 2002WO-US020025.  
 XX  
 PR 21-JUN-2001; 2001US-0299883P.  
 PR 23-APR-2002; 2002US-0375253P.  
 XX  
 PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
 XX  
 PI Fearon KL, Dina D, Tuck SF;

XX  
 DR WPI; 2003-210159/20.  
 XX  
 PT Novel chimeric immunomodulatory compound having immunomodulatory  
 PT activity, useful for modulating an immune response and for treating  
 PT cancer, has nucleic acid moieties and non-nucleic acid spacer moieties.  
 PS  
 PS Example 1; Page 110; 224pp; English.  
 CC The invention relates to a novel chimeric immunomodulatory compound (CIC)  
 CC having immunomodulatory activity, comprising two or more nucleic acid  
 CC moieties and one or more non-nucleic acid spacer moieties, where at least  
 CC one non-nucleic acid spacer moiety is covalently joined to two nucleic  
 CC acid moieties, where the spacer is not a polypeptide, and at least one  
 CC nucleic acid moiety comprises the sequence 5'-CG-3'. The chimeric  
 CC immunomodulatory compounds more specifically contain the nucleic acid  
 CC spacer moieties of linear hexaethylene glycol structure (HEG) subunits.  
 CC CIC's are useful for modulating an immune response in an individual,  
 CC where the individual suffers from a disorder associated with a Th2-type  
 CC immune response which is an allergy or allergy-induced asthma, and an  
 CC infectious disease. CIC is also useful for increasing IFN-gamma, or IFN-  
 CC alpha, in an individual, where the individual has idiopathic pulmonary  
 CC fibrosis, or a viral infection. CIC's are useful for ameliorating a  
 CC symptom of an infectious disease, or an immunoglobulin E (IgE)-related  
 CC disorder in an individual, where the IgE-related disorder is allergy, or  
 CC an allergy-related disorder. CIC's are also useful for treating cancer  
 CC and can be used for stimulating cellular immune system cells production  
 CC in an individual. This polynucleotide sequence represents a DNA sequence  
 CC of the invention.  
 XX  
 SQ Sequence 66 BP; 18 A; 9 C; 21 G; 18 T; 0 U; 0 Other;  
 Query Match 76.2%; Score 16; DB 8; Length 66;  
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 6 GAACGTTGAGATGAT 21  
 DB 8 GAACGTTGAGATGAT 23  
 RESULT 37  
 ID ABO75223 standard; DNA; 19 BP.  
 AC ABO75223;  
 XX  
 DT 05-NOV-2002 (first entry)  
 XX  
 DE ISS immunomodulatory oligonucleotide SEQ ID NO:58.  
 XX  
 XX Immunostimulatory sequence; ISS; immunomodulatory; immune response;  
 KW allergy; asthma; infectious disease; interferon-gamma; IFN-gamma;  
 KW idiopathic pulmonary fibrosis; viral infection; mycobacterial disease;  
 KW malaria; leishmaniasis; toxoplasmosis; schistosomiasis; clonorchiasis;  
 KW immunoglobulin E; IgE-related disorder; anti-allergic; antiasthmatic;  
 KW virucide; antibacterial; protozoacide; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO200252002-A2.  
 PD 04-JUL-2002.  
 XX  
 PF 27-DEC-2001; 2001WO-US050821.  
 XX  
 PR 27-DEC-2000; 2000US-0258675P.  
 XX  
 PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.  
 XX  
 PI Fearon KL, Dina D;

DR WPI; 2002-657426/70.

XX

PT Immunomodulatory polynucleotide for modulating an immune response in a

PT subject suffering from disorders associated with Th2-type immune

PT response, e.g. allergy, or infectious disease, comprises an

PT immunostimulatory sequence.

XX

PS Disclosure; Page 22; 95pp; English.

XX

CC The present invention describes an immunomodulatory polynucleotide (I)

CC comprising an immunostimulatory sequence (ISS). Also described: (1) an

CC immunomodulatory composition comprising (1); (2) an immunomodulatory

CC polynucleotide/microcarrier (IMP/MC) complex, comprising (1) linked to a

CC biodegradable MC, where the MC is less than 10 micrometre in size, and

CC (3) a kit comprising (1). (1) has anti-allergic, antiasthmatic, virucide,

CC antibacterial and protozoacidal activities, and can be used as a modulator

CC of immune response. (1) is useful for modulating an immune response in an

CC individual suffering from disorders associated with a Th2-type immune

CC response, especially an allergy or asthma, or an infectious disease. (1)

CC is also useful for increasing interferon-gamma (IFN-gamma) in an

CC individual having idiopathic pulmonary fibrosis, or IFN-alpha in an

CC individual having a viral infection. (1) is further useful for

CC ameliorating a symptom of an infectious disease caused by a cellular

CC pathogen such as mycobacterial disease, malaria, leishmaniasis,

CC toxoplasmosis, schistosomiasis and clonorchiasis in an individual, or a

CC symptom of an immunoglobulin E (IgE)-related disorder, preferably an

CC allergy-related disorder, in particular asthma in an individual. The

CC present sequence represents an immunomodulatory oligonucleotide from the

CC present invention

XX

SO Sequence 19 BP; 2 A; 4 C; 6 G; 7 T; 0 U; 0 Other;

Qy Query Match 75.2%; Score 15.8; DB 6; Length 19;

Best Local Similarity 89.5%; Pred. No. 1.5e+02;

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 1 TCGTCGAACGTTGAGATG 19

Db 1 TCGTCGTTGTTGAGATG 19

RESULT 38

AD888893

ID AD888893 standard; DNA; 19 BP.

XX

AC AD888893;

XX

DT 04-DEC-2003 (first entry)

XX

DE Chimeric immunomodulatory compound DNA sequence, SEQ ID No 96.

XX

KW chimeric immunomodulatory compound; CIC; immunomodulatory activity;

KW spacer moiety; linear hexaethylene glycol structure; HEG; immune;

KW Th2-type; allergy-induced asthma; infectious disease; IFN-gamma;

KW IFN-alpha; idiopathic pulmonary fibrosis; viral infection; ameliorating;

KW immunoglobulin E; IgE; allergy; cancer;

KW stimulating cellular immune system cell; ss.

XX

OS Synthetic.

XX

PN WO2003000922-A2.

XX

PD 03-JAN-2003.

XX

PF 21-JUN-2002; 2002WO-US020025.

XX

PR 21-JUN-2001; 2001US-0298883P.

XX

PR 23-APR-2002; 2002US-0375253P.

XX

PA (DYNA-) DYNAVAX TECHNOLOGIES CORP.

XX

PI Fearon KL, Dina D, Tuck SF;

XX

DR WPI; 2003-210159/20.

XX

PT Novel chimeric immunomodulatory compound having immunomodulatory

PT activity, useful for modulating an immune response and for treating

PT cancer, has nucleic acid moieties and non-nucleic acid spacer moieties.

XX

PS Disclosure; Page 35; 224pp; English.

XX

CC The invention relates to a novel chimeric immunomodulatory compound (CIC)

CC having immunomodulatory activity, comprising two or more nucleic acid

CC moieties and one or more non-nucleic acid spacer moieties, where at least

CC one non-nucleic acid spacer moiety is covalently joined to two nucleic

CC acid moieties, where the spacer is not a polypeptide, and at least one

CC nucleic acid moiety comprises the sequence 5'-CG-3'. The chimeric

CC immunomodulatory compounds more specifically contain the nucleic acid

CC spacer moieties of linear hexaethylene glycol structure (HEG) subunits.

CC CIC's are useful for modulating an immune response in an individual,

CC where the individual suffers from a disorder associated with a Th2-type

CC immune response which is an allergy or allergy-induced asthma, and an

CC infectious disease. CIC is also useful for increasing IFN-gamma, or IFN-

CC alpha, in an individual, where the individual has idiopathic pulmonary

CC fibrosis, or a viral infection. CIC's are useful for ameliorating a

CC symptom of an infectious disease, or an immunoglobulin E (IgE)-related

CC disorder in an individual, where the IgE-related disorder is allergy, or

CC an allergy-related disorder. CIC's are also useful for treating cancer

CC and can be used for stimulating cellular immune system cells production

CC in an individual. This polynucleotide sequence represents a DNA sequence

CC which is a nucleic acid moiety part of a chimeric immunomodulatory compound

CC of the invention.

XX

SO Sequence 19 BP; 2 A; 4 C; 6 G; 7 T; 0 U; 0 Other;

Qy Query Match 75.2%; Score 15.8; DB 8; Length 19;

Best Local Similarity 89.5%; Pred. No. 1.5e+02;

Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 1 TCGTCGAACGTTGAGATG 19

Db 1 TCGTCGTTGTTGAGATG 19

RESULT 39

ABL09441/C

ID ABL09441 standard; CDNA; 2726 BP.

XX

AC ABL09441;

XX

DT 26-MAR-2002 (first entry)

XX

DE Drosophila melanogaster expressed polynucleotide SEQ ID NO 22805.

XX

KW Drosophila; developmental biology; cell signalling; insecticide;

KW pharmaceutical; gene; ss.

XX

OS Drosophila melanogaster.

XX

PN WO200171042-A2.

XX

PD 27-SEP-2001.

XX

PF 23-MAR-2001; 2001WO-US009231.

XX

PR 23-MAR-2000; 2000US-0191637P.

XX

PR 11-JUL-2000; 2000US-00614150.

XX

PA (PEKE ) PE CORP NY.

XX

PI Venter JC, Adams M, Li PWD, Myers EW;

XX

DR WPI; 2001-656860/75.

XX

DR P-PSDB; ABB65338.

XX

PT New isolated nucleic acid detection reagent for detecting 1000 or more

PT genes from Drosophila and for elucidating cell signaling and cell-cell  
 PT interactions.

XX Claim 1; SEQ ID NO 22805; 21bp + Sequence Listing; English.

CC The invention relates to an isolated nucleic acid detection reagent  
 CC capable of detecting 1000 or more genes from Drosophila. The invention is  
 CC useful in developmental biology and in elucidating cell signaling and  
 CC cell-cell interactions in higher eukaryotes for the development of  
 CC insecticides, therapeutics and pharmaceutical drugs. The invention  
 CC discloses genomic DNA sequences (AB116176-AB130511), expressed DNA  
 CC sequences (AB101840-AB116175) and the encoded proteins (AB57737-  
 CC AB172072). The sequence data for this patent did not form part of the  
 CC printed specification, but was obtained in electronic format directly  
 CC from WIPO at ftp.wipo.int/pub/published\_pct\_sequences

XX Sequence 2726 BP; 659 A; 657 C; 781 G; 629 T; 0 U; 0 Other;

Query Match 75.2%; Score 15.8; DB 4; Length 2726;

Best Local Similarity 89.5%; Pred.No. 2.1e+02; Mismatches 2; Indels 0; Gaps 0;

DB 3 GTCGAACGTTGAGATGAT 21  
 155 GTCGAACGATCGTGATGAT 137

RESULT 40

AB109440/C  
 ID ABL09440 standard; cDNA; 5080 BP.

XX ABL09440;

DT 26-MAR-2002 (first entry)

DE Drosophila melanogaster expressed polynucleotide SEQ ID NO 22802.

XX Drosophila; developmental biology; cell signaling; insecticide;

KW pharmaceutical; gene; ss.

XX Drosophila melanogaster.

PN WO200171042-A2.

PD 27-SEP-2001.

PF 23-MAR-2001; 2001WO-US009231.

PR 23-MAR-2000; 2000US-0191637P.

PR 11-JUL-2000; 2000US-00614150.

PA (PEKE ) PE CORP NY.

PI Venter JC, Adams M, Li PWD, Myers EM;

DR WPI; 2001-656860/75.

DR P-PSDB; ABB65357.

PT New isolated nucleic acid detection reagent for detecting 1000 or more  
 PT genes from Drosophila and for elucidating cell signaling and cell-cell  
 PT interactions.

XX Claim 1; SEQ ID NO 22802; 21bp + Sequence Listing; English.

CC The invention relates to an isolated nucleic acid detection reagent  
 CC capable of detecting 1000 or more genes from Drosophila. The invention is  
 CC useful in developmental biology and in elucidating cell signaling and  
 CC cell-cell interactions in higher eukaryotes for the development of  
 CC insecticides, therapeutics and pharmaceutical drugs. The invention  
 CC discloses genomic DNA sequences (AB116176-AB130511), expressed DNA  
 CC sequences (AB101840-AB116175) and the encoded proteins (AB57737-  
 CC AB172072). The sequence data for this patent did not form part of the  
 CC printed specification, but was obtained in electronic format directly

CC from WIPO at ftp.wipo.int/pub/published\_pct\_sequences

XX Sequence 5080 BP; 1302 A; 1201 C; 1301 G; 1276 T; 0 U; 0 Other;

Query Match 75.2%; Score 15.8; DB 4; Length 5080;

Best Local Similarity 89.5%; Pred.No. 2.2e+02; Mismatches 2; Indels 0; Gaps 0;

DB 3 GTCGAACGTTGAGATGAT 21  
 1155 GTCGAACGATCGTGATGAT 1137

Search completed: April 6, 2004, 02:13:35  
 Job time : 401 secs



```

; APPLICANT: Van Nest, Gary
; APPLICANT: Tuck, Stephen
; APPLICANT: Fearon, Karen L.
; APPLICANT: Dina, Dino
; TITLE OF INVENTION: BIODEGRADABLE IMMUNOMODULATORY
; FILE REFERENCE: 37782001420
; CURRENT APPLICATION NUMBER: US/09/927,422A
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: U.S. 09/802,359
; PRIOR FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: U.S. 60/188,30
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 23
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-09-927-422A-16

```

```

Query Match          90.5%; Score 19; DB 10; Length 19;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```
QY      1 TCGTGAACGTTGAGATG 19
        |||||
DB      1 TCGTGAACGTTGAGATG 19

```

```

RESULT 3
US-10-033-243-19
; Sequence 19, Application US/10033243
; Publication No. US20030049266A1
; GENERAL INFORMATION:
; APPLICANT: FEARON, Karen L.
; APPLICANT: DINA, Dino
; TITLE OF INVENTION: IMMUNOMODULATORY POLYNUCLEOTIDES AND
; FILE REFERENCE: 37782001800
; CURRENT APPLICATION NUMBER: US/10/033,243
; PRIOR FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/258,675
; PRIOR FILING DATE: 2000-12-27
; NUMBER OF SEQ ID NOS: 133
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 19
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-10-033-243-19

```

```

Query Match          90.5%; Score 19; DB 14; Length 19;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```
QY      1 TCGTGAACGTTGAGATG 19
        |||||
DB      1 TCGTGAACGTTGAGATG 19

```

```

RESULT 4
US-10-176-883-41
; Sequence 41, Application US/10176883
; Publication No. US20030175731A1
; GENERAL INFORMATION:
; APPLICANT: FEARON, Karen
; APPLICANT: DINA, Dino
; APPLICANT: TUCK, Stephen

```

```

; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; FILE REFERENCE: 37782002000
; CURRENT APPLICATION NUMBER: US/10/176,883
; PRIOR FILING DATE: 2002-06-21
; PRIOR APPLICATION NUMBER: 60/299,883
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/375,253
; PRIOR FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 141
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 41
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
US-10-176-883-41

```

```

Query Match          90.5%; Score 19; DB 14; Length 19;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```
QY      1 TCGTGAACGTTGAGATG 19
        |||||
DB      1 TCGTGAACGTTGAGATG 19

```

```

RESULT 5
US-10-177-826-41
; Sequence 41, Application US/10177826
; Publication No. US20030199466A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; FILE REFERENCE: 37782002001
; CURRENT APPLICATION NUMBER: US/10/177,826
; PRIOR FILING DATE: 2002-06-21
; PRIOR APPLICATION NUMBER: 60/299,883
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/375,253
; PRIOR FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 141
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 41
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
US-10-177-826-41

```

```

Query Match          90.5%; Score 19; DB 14; Length 19;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```
QY      1 TCGTGAACGTTGAGATG 19
        |||||
DB      1 TCGTGAACGTTGAGATG 19

```

```

RESULT 6
US-10-033-243-30
; Sequence 30, Application US/10033243
; Publication No. US20030049266A1
; GENERAL INFORMATION:
; APPLICANT: FEARON, Karen L.
; APPLICANT: DINA, Dino
; TITLE OF INVENTION: IMMUNOMODULATORY POLYNUCLEOTIDES AND
; FILE REFERENCE: 37782001800
; CURRENT APPLICATION NUMBER: US/10/033,243
; PRIOR FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/258,675
; PRIOR FILING DATE: 2000-12-27
; NUMBER OF SEQ ID NOS: 133
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 19
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-10-033-243-30

```



```
FILE REFERENCE: 37782001800
; CURRENT APPLICATION NUMBER: US/10/033,243
; CURRENT FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/258,675
; PRIOR FILING DATE: 2000-12-27
; NUMBER OF SEQ ID NOS: 133
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 30
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-10-033-243-30
```

```
Query Match          90.5%; Score 19; DB 14; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.9;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY      1 TCGTGAACGTTTCGAGATG 19
         |||||
Db       4 TCGTGAACGTTTCGAGATG 22
```

```
RESULT 7
US-10-176-883-52
; Sequence 52, Application US/10176883
; Publication No. US20030175731A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; TITLE OF INVENTION: METHODS OF USING THE SAME-I
; FILE REFERENCE: 37782002000
; CURRENT APPLICATION NUMBER: US/10/176,883
; CURRENT FILING DATE: 2002-06-21
; PRIOR APPLICATION NUMBER: 60/299,883
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/375,253
; PRIOR FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 141
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 52
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
US-10-176-883-52
```

```
Query Match          90.5%; Score 19; DB 14; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.9;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY      1 TCGTGAACGTTTCGAGATG 19
         |||||
Db       4 TCGTGAACGTTTCGAGATG 22
```

```
RESULT 8
US-10-177-826-52
; Sequence 52, Application US/10177826
; Publication No. US20030199466A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; TITLE OF INVENTION: METHODS OF USING THE SAME-II
; FILE REFERENCE: 37782002001
; CURRENT APPLICATION NUMBER: US/10/177,826
; CURRENT FILING DATE: 2002-06-21
```

```
; PRIOR APPLICATION NUMBER: 60/299,883
; PRIOR FILING DATE: 2001-06-21,253
; PRIOR APPLICATION NUMBER: 60/375,253
; PRIOR FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 141
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 52
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
US-10-177-826-52
```

```
Query Match          90.5%; Score 19; DB 14; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.9;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY      1 TCGTGAACGTTTCGAGATG 19
         |||||
Db       4 TCGTGAACGTTTCGAGATG 22
```

```
RESULT 9
US-10-033-243-24
; Sequence 24, Application US/10033243
; Publication No. US20030049266A1
; GENERAL INFORMATION:
; APPLICANT: FEARON, Karen L.
; APPLICANT: DINA, Dino
; TITLE OF INVENTION: IMMUNOMODULATORY POLYNUCLEOTIDES AND
; TITLE OF INVENTION: METHODS OF USING THE SAME
; FILE REFERENCE: 37782001800
; CURRENT APPLICATION NUMBER: US/10/033,243
; CURRENT FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/258,675
; PRIOR FILING DATE: 2000-12-27
; NUMBER OF SEQ ID NOS: 133
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 24
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
; NAME/KEY: misc_feature
; LOCATION: 5
; OTHER INFORMATION: n = 5-bromocytosine
US-10-033-243-24
```

```
Query Match          85.7%; Score 18; DB 14; Length 19;
Best Local Similarity 94.7%; Pred. No. 13;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
OY      1 TCGTGAACGTTTCGAGATG 19
         |||||
Db       1 TCGTGAACGTTTCGAGATG 19
```

```
RESULT 10
US-10-176-883-46
; Sequence 46, Application US/10176883
; Publication No. US20030175731A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; TITLE OF INVENTION: METHODS OF USING THE SAME-I
; FILE REFERENCE: 37782002000
; CURRENT APPLICATION NUMBER: US/10/176,883
; CURRENT FILING DATE: 2002-06-21
```

```

; PRIOR APPLICATION NUMBER: 60/299,883
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/375,253
; PRIOR FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 141
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
; NAME/KEY: variation
; LOCATION: 5
; OTHER INFORMATION: n = 5-bromocytosine
US-10-176-883-46
```

```

Query Match      85.7%; Score 18; DB 14; Length 19;
Best Local Similarity 94.7%; Pred. No. 13;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY      1 TCGTCGAACGTTGAGATG 19
      |||||
DB      1 TCGTNGAACGTTGAGATG 19
```

```

RESULT 11
US-10-177-826-46
; Sequence 46, Application US/10177826
; Publication No. US20030199466A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; FILE REFERENCE: 377882002001
; CURRENT APPLICATION NUMBER: US/10/177,826
; CURRENT FILING DATE: 2002-06-21
; PRIOR APPLICATION NUMBER: 60/299,883
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/375,253
; PRIOR FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 141
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
; NAME/KEY: variation
; LOCATION: 5
; OTHER INFORMATION: n = 5-bromocytosine
US-10-177-826-46
```

```

Query Match      85.7%; Score 18; DB 14; Length 19;
Best Local Similarity 94.7%; Pred. No. 13;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY      1 TCGTCGAACGTTGAGATG 19
      |||||
DB      1 TCGTNGAACGTTGAGATG 19
```

```

RESULT 12
US-10-033-243-56
; Sequence 56, Application US/10033243
; Publication No. US20030049266A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen L.
```

```

; APPLICANT: DINA, Dino
; TITLE OF INVENTION: IMMUNOMODULATORY POLYNUCLEOTIDES AND
; FILE REFERENCE: 377882001800
; CURRENT APPLICATION NUMBER: US/10/033,243
; CURRENT FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/258,675
; PRIOR FILING DATE: 2000-12-27
; NUMBER OF SEQ ID NOS: 133
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 56
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-10-033-243-56
```

```

Query Match      82.9%; Score 17.4; DB 14; Length 19;
Best Local Similarity 94.7%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY      1 TCGTCGAACGTTGAGATG 19
      |||||
DB      1 TCGTCGACGTTGAGATG 19
```

```

RESULT 13
US-10-033-243-57
; Sequence 57, Application US/10033243
; Publication No. US20030049266A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen L.
; APPLICANT: DINA, Dino
; TITLE OF INVENTION: IMMUNOMODULATORY POLYNUCLEOTIDES AND
; FILE REFERENCE: 377882001800
; CURRENT APPLICATION NUMBER: US/10/033,243
; CURRENT FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/258,675
; PRIOR FILING DATE: 2000-12-27
; NUMBER OF SEQ ID NOS: 133
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 57
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-10-033-243-57
```

```

Query Match      82.9%; Score 17.4; DB 14; Length 19;
Best Local Similarity 94.7%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY      1 TCGTCGAACGTTGAGATG 19
      |||||
DB      1 TCGTCGACGTTGAGATG 19
```

```

RESULT 14
US-10-176-883-94
; Sequence 94, Application US/10176883
; Publication No. US20030175731A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; FILE REFERENCE: 377882002000
; CURRENT APPLICATION NUMBER: US/10/176,883
; CURRENT FILING DATE: 2002-06-21
```

; PRIOR APPLICATION NUMBER: 60/299,883  
; PRIOR FILING DATE: 2001-06-21  
; PRIOR APPLICATION NUMBER: 60/375,253  
; PRIOR FILING DATE: 2002-04-23  
; NUMBER OF SEQ ID NOS: 141  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 94  
; LENGTH: 19  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-10-176-883-94

Query Match 82.9%; Score 17.4; DB 14; Length 19;  
Best Local Similarity 94.7%; Pred. No. 26;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 TCGTGAACGTTGAGATG 19  
Db 1 TCGTGAACGTTGAGATG 19

RESULT 15  
US-10-176-883-95  
; Sequence 95, Application US/10176883  
; Publication No. US20030175731A1  
; GENERAL INFORMATION:  
; APPLICANT: Fearon, Karen  
; APPLICANT: Dina, Dino  
; APPLICANT: Tuck, Stephen  
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND  
; FILE REFERENCE: 377882002000  
; CURRENT FILING DATE: 2002-06-21  
; PRIOR APPLICATION NUMBER: 60/299,883  
; PRIOR FILING DATE: 2001-06-21  
; PRIOR APPLICATION NUMBER: 60/375,253  
; PRIOR FILING DATE: 2002-04-23  
; NUMBER OF SEQ ID NOS: 141  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 95  
; LENGTH: 19  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-10-176-883-95

Query Match 82.9%; Score 17.4; DB 14; Length 19;  
Best Local Similarity 94.7%; Pred. No. 26;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 TCGTGAACGTTGAGATG 19  
Db 1 TCGTGAACGTTGAGATG 19

RESULT 16  
US-10-177-826-94  
; Sequence 94, Application US/10177826  
; Publication No. US20030199466A1  
; GENERAL INFORMATION:  
; APPLICANT: Fearon, Karen  
; APPLICANT: Dina, Dino  
; APPLICANT: Tuck, Stephen  
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND  
; FILE REFERENCE: 377882002001  
; CURRENT APPLICATION NUMBER: US/10/177,826  
; CURRENT FILING DATE: 2002-06-21  
; PRIOR APPLICATION NUMBER: 60/299,883

; PRIOR FILING DATE: 2001-06-21  
; PRIOR APPLICATION NUMBER: 60/375,253  
; PRIOR FILING DATE: 2002-04-23  
; NUMBER OF SEQ ID NOS: 141  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 94  
; LENGTH: 19  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-10-177-826-94

Query Match 82.9%; Score 17.4; DB 14; Length 19;  
Best Local Similarity 94.7%; Pred. No. 26;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 TCGTGAACGTTGAGATG 19  
Db 1 TCGTGAACGTTGAGATG 19

RESULT 17  
US-10-177-826-95  
; Sequence 95, Application US/10177826  
; Publication No. US20030199466A1  
; GENERAL INFORMATION:  
; APPLICANT: Fearon, Karen  
; APPLICANT: Dina, Dino  
; APPLICANT: Tuck, Stephen  
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND  
; FILE REFERENCE: 377882002001  
; CURRENT FILING DATE: 2002-06-21  
; PRIOR APPLICATION NUMBER: 60/299,883  
; PRIOR FILING DATE: 2001-06-21  
; PRIOR APPLICATION NUMBER: 60/375,253  
; PRIOR FILING DATE: 2002-04-23  
; NUMBER OF SEQ ID NOS: 141  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 95  
; LENGTH: 19  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-10-177-826-95

Query Match 82.9%; Score 17.4; DB 14; Length 19;  
Best Local Similarity 94.7%; Pred. No. 26;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 TCGTGAACGTTGAGATG 19  
Db 1 TCGTGAACGTTGAGATG 19

RESULT 18  
US-10-033-243-23  
; Sequence 23, Application US/10033243  
; Publication No. US20030049266A1  
; GENERAL INFORMATION:  
; APPLICANT: FEARON, Karen L.  
; APPLICANT: DINA, Dino  
; TITLE OF INVENTION: IMMUNOMODULATORY POLYNUCLEOTIDES AND  
; FILE REFERENCE: 377882001800  
; CURRENT APPLICATION NUMBER: US/10/033,243  
; CURRENT FILING DATE: 2002-04-03  
; PRIOR APPLICATION NUMBER: 60/258,675  
; PRIOR FILING DATE: 2000-12-27  
; NUMBER OF SEQ ID NOS: 133

```
SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 23
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
; NAME/KEY: misc_feature
; LOCATION: 2..5
; OTHER INFORMATION: n = 5-bromocytosine
US-10-033-243-23

Query Match      81.0%; Score 17; DB 14; Length 19;
Best Local Similarity 89.5%; Pred. No. 42;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGACGTTGAGATG 19
   |||||
Db 1 TNGTNGACGTTGAGATG 19

RESULT 19
US-10-176-883-45
; Sequence 45, Application US/10176883
; Publication No. US20030175731A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; FILE REFERENCE: 377882002000
; CURRENT FILING DATE: 2002-06-21
; PRIOR FILING DATE: 2001-06-21
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/375,253
; PRIOR FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 141
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 45
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
; NAME/KEY: variation
; LOCATION: 2..5
; OTHER INFORMATION: n = 5-bromocytosine
US-10-176-883-45

Query Match      81.0%; Score 17; DB 14; Length 19;
Best Local Similarity 89.5%; Pred. No. 42;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGACGTTGAGATG 19
   |||||
Db 1 TNGTNGACGTTGAGATG 19

RESULT 20
US-10-177-826-45
; Sequence 45, Application US/10177826
; Publication No. US20030199466A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; FILE REFERENCE: 377882002001
```

```
CURRENT APPLICATION NUMBER: US/10/177,826
; CURRENT FILING DATE: 2002-06-21
; PRIOR APPLICATION NUMBER: 60/299,883
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/375,253
; PRIOR FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 141
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 45
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic construct
; NAME/KEY: variation
; LOCATION: 2..5
; OTHER INFORMATION: n = 5-bromocytosine
US-10-177-826-45

Query Match      81.0%; Score 17; DB 14; Length 19;
Best Local Similarity 89.5%; Pred. No. 42;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGACGTTGAGATG 19
   |||||
Db 1 TNGTNGACGTTGAGATG 19

RESULT 21
US-10-033-243-14
; Sequence 14, Application US/10033243
; Publication No. US20030049266A1
; GENERAL INFORMATION:
; APPLICANT: FEARON, Karen L.
; APPLICANT: DINA, Dino
; TITLE OF INVENTION: IMMUNOMODULATORY POLYNUCLEOTIDES AND
; FILE REFERENCE: 377882001800
; CURRENT APPLICATION NUMBER: US/10/033,243
; CURRENT FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/258,675
; PRIOR FILING DATE: 2000-12-27
; NUMBER OF SEQ ID NOS: 133
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Polynucleotide containing CG
US-10-033-243-14

Query Match      78.1%; Score 16.4; DB 14; Length 18;
Best Local Similarity 94.4%; Pred. No. 87;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 CGTCGACGTTGAGATG 19
   |||||
Db 1 CTTGCAACGTTGAGATG 18

RESULT 22
US-10-176-883-36
; Sequence 36, Application US/10176883
; Publication No. US20030175731A1
; GENERAL INFORMATION:
; APPLICANT: Fearon, Karen
; APPLICANT: Dina, Dino
; APPLICANT: Tuck, Stephen
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND
; FILE REFERENCE: 377882002000
```

```

RESULT 26
US-09-802-376-3/c
: Sequence 3, Application US/09802376
: Patent No. US20020055477A1
: GENERAL INFORMATION:
: APPLICANT: Van Nest, Gary
: APPLICANT: Tuck, Stephen
: TITLE OF INVENTION: IMMUNOMODULATORY FORMULATIONS AND METHODS FOR USE THEREOF
: FILE REFERENCE: 37788201700
: CURRENT APPLICATION NUMBER: US/09/802,376
: CURRENT FILING DATE: 2001-03-09
: PRIOR APPLICATION NUMBER: 60/188,557
: PRIOR FILING DATE: 2000-03-10
: NUMBER OF SEQ. ID NOS.: 11
: SOFTWARE: FastSeq for Windows Version 4.0
: SEQ ID NO 3
: LENGTH: 23
: TYPE: DNA

```

ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Polynucleotide containing CG  
US-09-802-376-3

Query Match 78.1%; Score 16.4; DB 9; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GTCGAACGTTGAGATGA 20  
|||  
18 GTGGAACGTTGAGATGA 1

DB 18 GTGGAACGTTGAGATGA 1

RESULT 27  
US-09-802-370-3/c

Sequence 3, Application US/09802370  
Patent No. US20020098199A1

GENERAL INFORMATION:

APPLICANT: Van Nest, Gary

APPLICANT: Eiden, Joseph J. Jr.

TITLE OF INVENTION: METHODS OF SUPPRESSING HEPATITIS VIRUS

TITLE OF INVENTION: INFECTION USING IMMUNOMODULATORY POLYNUCLEOTIDE SEQUENCES

FILE REFERENCE: 377882001200

CURRENT APPLICATION NUMBER: US/09/802,370

PRIOR FILING DATE: 2001-09-24

PRIOR APPLICATION NUMBER: 60/188,301

NUMBER OF SEQ ID NOS: 8

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 3

LENGTH: 23

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE:

OTHER INFORMATION: Polynucleotide containing CG

US-09-802-370-3

Query Match 78.1%; Score 16.4; DB 9; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GTCGAACGTTGAGATGA 20  
|||  
18 GTGGAACGTTGAGATGA 1

DB 18 GTGGAACGTTGAGATGA 1

RESULT 28  
US-09-802-445-3/c

Sequence 3, Application US/09802445  
Patent No. US20020107212A1

GENERAL INFORMATION:

APPLICANT: Van Nest, Gary

APPLICANT: Eiden, Joseph J. Jr.

TITLE OF INVENTION: METHODS OF REDUCING PAPILLOMAVIRUS INFECTION USING IMMUNOMODULATORY

FILE REFERENCE: 377882001300

CURRENT APPLICATION NUMBER: US/09/802,445

PRIOR FILING DATE: 2001-09-24

PRIOR APPLICATION NUMBER: 60/188,265

NUMBER OF SEQ ID NOS: 8

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 3

LENGTH: 23

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE:

OTHER INFORMATION: Polynucleotide containing CG

US-09-802-445-3

Query Match 78.1%; Score 16.4; DB 9; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GTCGAACGTTGAGATGA 20  
|||  
18 GTGGAACGTTGAGATGA 1

DB 18 GTGGAACGTTGAGATGA 1

RESULT 29  
US-09-927-422A-3/c

Sequence 3, Application US/09927422A  
Publication No. US20030022852A1

GENERAL INFORMATION:

APPLICANT: Van Nest, Gary

APPLICANT: Tuck, Stephen

APPLICANT: Fearon, Karen L.

APPLICANT: Dina, Dino

TITLE OF INVENTION: BIODEGRADABLE IMMUNOMODULATORY

TITLE OF INVENTION: FORMULATIONS AND METHODS FOR USE THEREOF

FILE REFERENCE: 377882001420

CURRENT APPLICATION NUMBER: US/09/927,422A

PRIOR FILING DATE: 2001-08-10

PRIOR APPLICATION NUMBER: U.S. 09/802,359

PRIOR FILING DATE: 2001-03-09

PRIOR APPLICATION NUMBER: U.S. 60/188,301

NUMBER OF SEQ ID NOS: 23

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 3

LENGTH: 23

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE:

OTHER INFORMATION: Polynucleotide containing CG

US-09-927-422A-3

Query Match 78.1%; Score 16.4; DB 10; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GTCGAACGTTGAGATGA 20  
|||  
18 GTGGAACGTTGAGATGA 1

DB 18 GTGGAACGTTGAGATGA 1

RESULT 30  
US-09-927-884-3/c

Sequence 3, Application US/09927884  
Publication No. US20030059773A1

GENERAL INFORMATION:

APPLICANT: Van Nest, Gary

APPLICANT: Tuck, Stephen

APPLICANT: Fearon, Karen L.

APPLICANT: Dina, Dino

TITLE OF INVENTION: IMMUNOMODULATORY FORMULATIONS AND

FILE REFERENCE: 377882001720

CURRENT APPLICATION NUMBER: US/09/927,884

PRIOR FILING DATE: 2001-08-10

PRIOR APPLICATION NUMBER: U.S. 09/802,376

PRIOR FILING DATE: 2001-03-09

PRIOR APPLICATION NUMBER: U.S. 60/188,557

NUMBER OF SEQ ID NOS: 14

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 3

LENGTH: 23

TYPE: DNA

ORGANISM: Artificial Sequence

FEATURE:

OTHER INFORMATION: Polynucleotide containing CG

US-09-927-884-3

Query Match 78.1%; Score 16.4; DB 10; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;

Best Local Similarity 94.4%; Pred. No. 88;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 3 GTCGAACGTCGAGATGA 20  
|||  
Db 18 GTGGAACGTCGAGATGA 1

## RESULT 31

US-09-802-359-3/c  
; Sequence 3, Application US/09802359  
; Publication No. US20030129251A1  
; GENERAL INFORMATION:  
; APPLICANT: Van Nest, Gary  
; APPLICANT: Tuck, Stephen  
; TITLE OF INVENTION: IMMUNOMODULATORY FORMULATIONS AND METHODS FOR USE THEREOF  
; FILE REFERENCE: 37788201400  
; CURRENT APPLICATION NUMBER: US/09/802,359  
; PRIOR FILING DATE: 2001-03-09  
; PRIOR APPLICATION NUMBER: 60/188,303  
; PRIOR FILING DATE: 2000-03-10  
; NUMBER OF SEQ ID NOS: 11  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 3  
; LENGTH: 23  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Polynucleotide containing CG  
US-09-802-359-3

Query Match 78.1%; Score 16.4; DB 10; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 3 GTCGAACGTCGAGATGA 20  
|||  
Db 18 GTGGAACGTCGAGATGA 1

## RESULT 32

US-10-357-760-3/c  
; Sequence 3, Application US/10357760  
; Publication No. US20030216340A1  
; GENERAL INFORMATION:  
; APPLICANT: Van Nest, Gary  
; APPLICANT: Eiden, Joseph J. Jr.  
; TITLE OF INVENTION: METHODS OF SUPPRESSING HEPATITIS VIRUS  
; TITLE OF INVENTION: INFECTION USING IMMUNOMODULATORY POLYNUCLEOTIDE SEQUENCES  
; FILE REFERENCE: 377882001200  
; CURRENT APPLICATION NUMBER: US/10/357,760  
; CURRENT FILING DATE: 2003-02-03  
; PRIOR APPLICATION NUMBER: US/09/802,370  
; PRIOR FILING DATE: 2001-09-24  
; PRIOR APPLICATION NUMBER: 60/188,301  
; PRIOR FILING DATE: 2000-03-10  
; NUMBER OF SEQ ID NOS: 8  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 3  
; LENGTH: 23  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Polynucleotide containing CG  
US-10-357-760-3

Query Match 78.1%; Score 16.4; DB 15; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 3 GTCGAACGTCGAGATGA 20  
|||  
Db 18 GTGGAACGTCGAGATGA 1

RESULT 33  
US-10-413-504-4/c  
; Sequence 4, Application US/10413504  
; Publication No. US20040006034A1  
; GENERAL INFORMATION:  
; APPLICANT: Raz, Eyal  
; APPLICANT: Schwartz, David  
; APPLICANT: Roman, Mark  
; APPLICANT: Dina, Dino  
; TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,  
; TITLE OF INVENTION: COMPOSITIONS THEREOF AND METHODS OF USE THEREOF  
; FILE REFERENCE: 37788-20004.01  
; CURRENT APPLICATION NUMBER: US/10/413,504  
; CURRENT FILING DATE: 2003-04-11  
; PRIOR APPLICATION NUMBER: US 09/296,477  
; PRIOR FILING DATE: 1999-04-22  
; PRIOR APPLICATION NUMBER: US 09/092,329  
; PRIOR FILING DATE: 1998-06-05  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 4  
; LENGTH: 23  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Immunostimulatory oligonucleotide  
US-10-413-504-4

Query Match 78.1%; Score 16.4; DB 15; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 3 GTCGAACGTCGAGATGA 20  
|||  
Db 18 GTGGAACGTCGAGATGA 1

## RESULT 34

US-10-426-237-3/c  
; Sequence 3, Application US/10426237  
; Publication No. US20040009942A1  
; GENERAL INFORMATION:  
; APPLICANT: Dynavax Technologies Corporation  
; APPLICANT: Van Nest, Gary  
; TITLE OF INVENTION: METHODS OF PREVENTING AND TREATING  
; TITLE OF INVENTION: RESPIRATORY VIRAL INFECTION USING IMMUNOMODULATORY  
; TITLE OF INVENTION: POLYNUCLEOTIDE SEQUENCES  
; FILE REFERENCE: 377882000920  
; CURRENT APPLICATION NUMBER: US/10/426,237  
; CURRENT FILING DATE: 2003-04-29  
; PRIOR APPLICATION NUMBER: 09/802,686  
; PRIOR FILING DATE: 2001-03-09  
; PRIOR APPLICATION NUMBER: 60/188,583  
; PRIOR FILING DATE: 2000-03-10  
; NUMBER OF SEQ ID NOS: 10  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 3  
; LENGTH: 23  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Polynucleotide containing CG  
US-10-426-237-3

Query Match 78.1%; Score 16.4; DB 15; Length 23;  
Best Local Similarity 94.4%; Pred. No. 88;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 3 GTCGAACGTCGAGATGA 20  
|||  
Db 18 GTGGAACGTCGAGATGA 1

```
RESULT 35
US-10-424-599-21073/c
; Sequence 21073, Application US/10424599
; Publication No. US20040031072A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J
; APPLICANT: Kovalic, David K
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Soy Nucleic Acid Molecules and Other Molecules Associated With
; TITLE OF INVENTION: Plants and Uses Thereof for Plant Improvement
; FILE REFERENCE: 38-21(53223)B
; CURRENT APPLICATION NUMBER: US/10/424,599
; NUMBER OF SEQ ID NOS: 285684
; SEQ ID NO 21073
; LENGTH: 450
; TYPE: DNA
; ORGANISM: Glycine max
; FEATURE:
; OTHER INFORMATION: Clone ID: PAT_MRT3847_119033C.1
US-10-424-599-21073

Query Match      78.1%; Score 16.4; DB 12; Length 450;
Best Local Similarity 94.4%; Pred. No. 1.1e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      4 TCGAACGTTGAGATGAT 21
      |||||
Db      373 TCGAAGTTTCGAGATGAT 356

RESULT 36
US-10-282-122A-35541
; Sequence 35541, Application US/10282122A
; Publication No. US20040029129A1
; GENERAL INFORMATION:
; APPLICANT: Wang, Liangsu
; APPLICANT: Zamudio, Carlos
; APPLICANT: Malone, Cheryl
; APPLICANT: Haselbeck, Robert
; APPLICANT: Ohlsen, Kari
; APPLICANT: Zyskind, Judith
; APPLICANT: Wall, Daniel
; APPLICANT: Trawick, John
; APPLICANT: Carr, Grant
; APPLICANT: Yamamoto, Robert
; APPLICANT: Forsyth, R.
; APPLICANT: Xu, H.
; TITLE OF INVENTION: Identification of Essential Genes in Microorganisms
; FILE REFERENCE: ELITRA.034A
; CURRENT APPLICATION NUMBER: US/10/282,122A
; CURRENT FILING DATE: 2003-02-20
; PRIOR APPLICATION NUMBER: 60/191,078
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: 60/206,848
; PRIOR FILING DATE: 2000-05-23
; PRIOR APPLICATION NUMBER: 60/207,727
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: 60/230,335
; PRIOR FILING DATE: 2000-09-06
; PRIOR APPLICATION NUMBER: 60/230,347
; PRIOR FILING DATE: 2000-09-09
; PRIOR APPLICATION NUMBER: 60/242,578
; PRIOR FILING DATE: 2000-10-23
; PRIOR APPLICATION NUMBER: 60/257,931
; PRIOR FILING DATE: 2000-11-27
; PRIOR APPLICATION NUMBER: 60/267,636
; PRIOR FILING DATE: 2001-02-09
```

```
; PRIOR APPLICATION NUMBER: 60/269,308
; PRIOR FILING DATE: 2001-02-16
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 78614
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 35541
; LENGTH: 1020
; TYPE: DNA
; ORGANISM: Staphylococcus haemolyticus
US-10-282-122A-35541

Query Match      77.1%; Score 16.2; DB 12; Length 1020;
Best Local Similarity 85.7%; Pred. No. 1.4e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy      1 TCGTGACGTTGAGATGAT 21
      |||||
Db      922 TTGTGAATGTTGAGATGAT 942

RESULT 37
US-10-282-122A-21830/c
; Sequence 21830, Application US/10282122A
; Publication No. US20040029129A1
; GENERAL INFORMATION:
; APPLICANT: Wang, Liangsu
; APPLICANT: Zamudio, Carlos
; APPLICANT: Malone, Cheryl
; APPLICANT: Haselbeck, Robert
; APPLICANT: Ohlsen, Kari
; APPLICANT: Zyskind, Judith
; APPLICANT: Wall, Daniel
; APPLICANT: Trawick, John
; APPLICANT: Carr, Grant
; APPLICANT: Yamamoto, Robert
; APPLICANT: Forsyth, R.
; APPLICANT: Xu, H.
; TITLE OF INVENTION: Identification of Essential Genes in Microorganisms
; FILE REFERENCE: ELITRA.034A
; CURRENT APPLICATION NUMBER: US/10/282,122A
; CURRENT FILING DATE: 2003-02-20
; PRIOR APPLICATION NUMBER: 60/191,078
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: 60/206,848
; PRIOR FILING DATE: 2000-05-23
; PRIOR APPLICATION NUMBER: 60/207,727
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: 60/230,335
; PRIOR FILING DATE: 2000-09-06
; PRIOR APPLICATION NUMBER: 60/230,347
; PRIOR FILING DATE: 2000-09-09
; PRIOR APPLICATION NUMBER: 60/242,578
; PRIOR FILING DATE: 2000-10-23
; PRIOR APPLICATION NUMBER: 60/253,625
; PRIOR FILING DATE: 2000-11-27
; PRIOR APPLICATION NUMBER: 60/257,931
; PRIOR FILING DATE: 2000-12-22
; PRIOR APPLICATION NUMBER: 60/267,636
; PRIOR FILING DATE: 2001-02-09
; PRIOR APPLICATION NUMBER: 60/269,308
; PRIOR FILING DATE: 2001-02-16
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 78614
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 21830
; LENGTH: 3126
; TYPE: DNA
; ORGANISM: Enterococcus faecium
US-10-282-122A-21830

Query Match      77.1%; Score 16.2; DB 12; Length 3126;
Best Local Similarity 85.7%; Pred. No. 1.5e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
```



OY 1 TCGTCGAACGTTCCGAGATGAT 21  
|||||  
Db 2719 TCGTTAAACGTTGCGAGATGAT 2699

## RESULT 38

US-10-424-599-28340/C  
; Sequence 28340, Application US/10424599  
; Publication No. US20040031072A1  
; GENERAL INFORMATION:  
; APPLICANT: La Rosa Thomas J  
; APPLICANT: Kovalic David K  
; APPLICANT: Zhou Yihua  
; APPLICANT: Cao Yongwei  
; TITLE OF INVENTION: Soy Nucleic Acid Molecules and Other Molecules Associated with  
; TITLE OF INVENTION: Plants and Uses Thereof for Plant Improvement  
; FILE REFERENCE: 38-21(53223) B  
; CURRENT APPLICATION NUMBER: US/10-424,599  
; CURRENT FILING DATE: 2003-04-28  
; NUMBER OF SEQ ID NOS: 285684  
; SEQ ID NO 28340  
; LENGTH: 3999  
; TYPE: DNA  
; ORGANISM: Glycine max  
; FEATURE:  
; OTHER INFORMATION: Clone ID: PAT\_MRT3847\_125592C.1  
US-10-424-599-28340

Query Match 77.1%; Score 16.2; DB 12; Length 3999;  
Best Local Similarity 85.7%; Pred. No. 1.6e+02;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 TCGTCGAACGTTCCGAGATGAT 21  
|||||  
Db 1863 TCGTCGAACGTTCCGAGATGAT 1843

## RESULT 39

US-10-176-883-139  
; Sequence 139, Application US/10176883  
; Publication No. US2003015731A1  
; GENERAL INFORMATION:  
; APPLICANT: Fearon, Karen  
; APPLICANT: Dina, Dino  
; APPLICANT: Tuck, Stephen  
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND  
; TITLE OF INVENTION: METHODS OF USING THE SAME-I  
; FILE REFERENCE: 377882002000  
; CURRENT APPLICATION NUMBER: US/10-176,883  
; CURRENT FILING DATE: 2002-06-21  
; PRIOR APPLICATION NUMBER: 60/299,883  
; PRIOR FILING DATE: 2001-06-21  
; PRIOR APPLICATION NUMBER: 60/375,253  
; PRIOR FILING DATE: 2002-04-23  
; NUMBER OF SEQ ID NOS: 141  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 139  
; LENGTH: 66  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-10-176-883-139

Query Match 76.2%; Score 16; DB 14; Length 66;  
Best Local Similarity 100.0%; Pred. No. 1.5e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 GAACGTTCCGAGATGAT 21  
|||||  
Db 8 GAACGTTCCGAGATGAT 23

RESULT 40  
US-10-177-826-139  
; Sequence 139, Application US/10177826  
; Publication No. US20030199466A1  
; GENERAL INFORMATION:  
; APPLICANT: Fearon, Karen  
; APPLICANT: Dina, Dino  
; APPLICANT: Tuck, Stephen  
; TITLE OF INVENTION: CHIMERIC IMMUNOMODULATORY COMPOUNDS AND  
; TITLE OF INVENTION: METHODS OF USING THE SAME-II  
; FILE REFERENCE: 377882002001  
; CURRENT APPLICATION NUMBER: US/10-177,826  
; CURRENT FILING DATE: 2002-06-21  
; PRIOR APPLICATION NUMBER: 60/299,883  
; PRIOR FILING DATE: 2001-06-21  
; PRIOR APPLICATION NUMBER: 60/375,253  
; PRIOR FILING DATE: 2002-04-23  
; NUMBER OF SEQ ID NOS: 141  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 139  
; LENGTH: 66  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-10-177-826-139

Query Match 76.2%; Score 16; DB 14; Length 66;  
Best Local Similarity 100.0%; Pred. No. 1.5e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 GAACGTTCCGAGATGAT 21  
|||||  
Db 8 GAACGTTCCGAGATGAT 23

Search completed: April 6, 2004, 04:04:03  
Job time : 346 secs

**This Page Blank (uspto)**

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## OM nucleic - nucleic search, using SW model

Run on: April 6, 2004, 01:41:36 ; Search time 75 Seconds

(without alignments)  
155.386 Million cell updates/sec

Title: US-10-033-243-132

Perfect score: 21

Sequence: 1 cgcgcgaacgttcgagatgat 21

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 682709 seqs, 277475446 residues

Total number of hits satisfying chosen parameters: 1365418

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA.\*  
1: /cgn2\_6/ptodata/2/ina/5A.COMB.seq:\*  
2: /cgn2\_6/ptodata/2/ina/5B.COMB.seq:\*  
3: /cgn2\_6/ptodata/2/ina/6A.COMB.seq:\*  
4: /cgn2\_6/ptodata/2/ina/6B.COMB.seq:\*  
5: /cgn2\_6/ptodata/2/ina/PCBUS.COMB.seq:\*  
6: /cgn2\_6/ptodata/2/ina/backfilese1.seq:\*

Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16.4	78.1	23	4	US-09-296-477-4
2	16.2	77.1	3135	4	US-09-107-532A-1575
3	15.4	73.3	816	3	US-08-776-251-10
4	15.4	73.3	816	3	US-08-776-251-10
5	15.4	73.3	1900	4	US-09-555-000-1
6	15.2	72.4	342	4	US-09-134-000C-2551
7	15.2	72.4	4590	4	US-09-134-001C-1108
8	15.2	72.4	10619	4	US-10-204-708-4
9	15	71.4	22	4	US-09-235-742-19
10	15	71.4	22	4	US-09-347-343-32
11	15	71.4	22	4	US-09-820-484-1
12	15	71.4	22	4	US-09-820-484-3
13	15	71.4	22	4	US-09-774-403A-1
14	15	71.4	22	4	US-09-296-477-1
15	15	71.4	22	4	US-09-296-477-2
16	15	71.4	22	4	US-09-296-477-5
17	15	71.4	22	4	US-09-308-036A-1
18	15	71.4	22	4	US-09-791-500-1
19	14.8	70.5	813	4	US-09-107-532A-1566
20	14.8	70.5	1069	4	US-09-374-174B-1
21	14.8	70.5	1497	4	US-09-252-991A-2256
22	14.8	70.5	1950	4	US-08-252-991A-2425
23	14.8	70.5	2799	1	US-08-446-794A-5
24	14.6	69.5	424	4	US-09-621-976-12664
25	14.6	69.5	3468	4	US-09-221-017B-893
26	14.6	69.5	77	1	US-08-399-412A-58
27	14.4	68.6	77	1	US-08-399-412A-58

## ALIGNMENTS

28	14.2	67.6	77	1	US-08-447-169A-36	Sequence 36, Appl
29	14.2	67.6	77	2	US-08-233-012C-36	Sequence 36, Appl
30	14.2	67.6	1535	4	US-09-848-295-1	Sequence 1, Appl
31	14.2	67.6	4715	4	US-08-956-171E-203	Sequence 203, App
32	14.2	67.6	14636	3	US-09-173-914-6	Sequence 6, Appl
33	14.2	67.6	50937	3	US-09-428-517-1	Sequence 1, Appl
34	14.2	67.6	4403765	3	US-09-103-840A-2	Sequence 2, Appl
35	14.2	67.6	4411529	3	US-09-103-840A-1	Sequence 1, Appl
36	14	66.7	22	4	US-09-296-477-15	Sequence 15, Appl
37	13.8	65.7	66	2	US-08-710-134-63	Sequence 63, Appl
38	13.8	65.7	66	2	US-08-485-885-63	Sequence 63, Appl
39	13.8	65.7	86	2	US-08-710-134-64	Sequence 64, Appl
40	13.8	65.7	86	2	US-08-485-885-64	Sequence 64, Appl
41	13.8	65.7	173	4	US-09-313-294A-3045	Sequence 3045, Ap
42	13.8	65.7	223	4	US-08-956-171E-1271	Sequence 1271, Ap
43	13.8	65.7	321	3	US-09-240-274-197	Sequence 197, App
44	13.8	65.7	459	4	US-09-540-236-1391	Sequence 1391, Ap
45	13.8	65.7	531	4	US-09-134-000C-1500	Sequence 1500, Ap

RESULT 1  
US-09-296-477-4/C  
Sequence 4, Application US/09296477A  
Patent No. 6589940  
GENERAL INFORMATION:  
APPLICANT: RAZ, E.  
APPLICANT: SCHWARTZ, D.  
APPLICANT: ROMAN, M.  
APPLICANT: DIMA, D.  
TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,  
TITLE OF INVENTION: COMPOSITIONS THEREOF AND METHODS OF USE  
FILE REFERENCE: 377882000420  
CURRENT APPLICATION NUMBER: US/09/296,477A  
CURRENT FILING DATE: 1999-04-22  
EARLIER APPLICATION NUMBER: 09/092,329  
EARLIER FILING DATE: 1998-06-05  
EARLIER APPLICATION NUMBER: 60/048,793  
EARLIER FILING DATE: 1997-06-06  
NUMBER OF SEQ ID NOS: 21  
SOFTWARE: FASTSEQ for Windows Version 3.0  
SEQ ID NO 4  
LENGTH: 23  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Synthetic construct  
US-09-296-477-4  
Query Match 78.1%; Score 16.4; DB 4; Length 23;  
Best Local Similarity 94.4%; Pred No. 5.5; Indels 0; Gaps 0;  
Matches 17; Conservative 0; Mismatches 1;  
Db 3 GTGGAACGTCGAGATGA 20  
18 GTGGAACGTCGAGATGA 1  
RESULT 2  
US-09-107-532A-1575/C  
Sequence 1575, Application US/09107532A  
Patent No. 6583275  
GENERAL INFORMATION:  
APPLICANT: Lynn A Doucette-Stamm and David Bush  
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO  
ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS  
NUMBER OF SEQUENCES: 7310  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: GENOME THERAPEUTICS CORPORATION  
STREET: 100 Beaver Street

```

CITY: Waltham
STATE: Massachusetts
COUNTRY: USA
ZIP: 02354

COMPUTER READABLE FORM:
MEDIUM TYPE: CD/ROM ISO9660
COMPUTER: PC
OPERATING SYSTEM: <Unknown>
SOFTWARE: ASCII

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/107,532A
FILING DATE: 30-Jun-1998

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/085,598
FILING DATE: 14 May 1998
APPLICATION NUMBER: 60/051571
FILING DATE: July 2, 1997

ATTORNEY/AGENT INFORMATION:
NAME: Ariniello, Pamela Deneke
REGISTRATION NUMBER: 40,489
REFERENCE/DOCKET NUMBER: GTC-012
TELECOMMUNICATION INFORMATION:
TELEPHONE: (781)893-5007
TELEFAX: (781)893-8277

INFORMATION FOR SEQ ID NO: 1575:
SEQUENCE CHARACTERISTICS:
LENGTH: 3135 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: circular
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Enterococcus faecium

FEATURE:
NAME/KEY: misc_feature
LOCATION: (8) LOCATION 1..3135
SEQUENCE DESCRIPTION: SEQ ID NO: 1575:
US-09-107-532A-1575

Query Match      77.1%; Score 16.2; DB 4; Length 3135;
Beac local Similarity 85.7%; Pred. No. 13;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Oy      1 TCGTCGACGTTTCGAGATGAT 21
      ||||| ||||| ||||| |||||
Db      2725 TCGTTAACGTTTCGAGATGAT 2705

RESULT 3
US-08-776-251-10
Sequence 10, Application US/08776251
Patent No. 6025340
GENERAL INFORMATION:
APPLICANT: Springer, Caroline J
APPLICANT: Marais, Richard
TITLE OF INVENTION: Surface expression of enzyme in gene directed prodrug therapy
NUMBER OF SEQUENCES: 27
CORRESPONDENCE ADDRESS:
ADDRESSEE: Nixon & Vanderhye
STREET: 1100 No. 6025340th Glebe Road, 8th Floor
CITY: Arlington
STATE: Virginia
COUNTRY: USA

COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/776,251
FILING DATE: 31-JAN-1997

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      PRIOR APPLICATION DATA:
      APPLICATION NUMBER: PCT/GB95/01782
      FILING DATE: 27-JUL-1995
      PRIOR APPLICATION DATA:
      APPLICATION NUMBER: GB 9415167.7
      FILING DATE: 27-JUL-1994
      ATTORNEY/AGENT INFORMATION:
      NAME: Arthur R. Crawford
      REGISTRATION NUMBER: 25,327
      REFERENCE/DOCKET NUMBER: 620-20
      INFORMATION FOR SEQ ID NO: 10:
      SEQUENCE CHARACTERISTICS:
      LENGTH: 816 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
      MOLECULE TYPE: CDNA
      US-08-776-251-10

      Query Match      73.3%; Score 15.4; DB 3; Length 816;
      Best Local Similarity 94.1%; Pred. No. 31;
      Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

      4 TCGAACGTTGAGATGA 20
      |||
      Db      619 TCGAACGTTGAGACGA 635

      RESULT 4
      US-08-776-251-10/c
      Sequence 10. Application US/08776251
      Patent No. 6025340
      GENERAL INFORMATION:
      APPLICANT: Springer, Caroline J
      APPLICANT: Marais, Richard
      TITLE OF INVENTION: Surface expression of enzyme in gene directed prodrug therapy
      NUMBER OF SEQUENCES: 27
      CORRESPONDENCE ADDRESS:
      ADDRESSEE: Nixon & Vanderhye
      STREET: 1100 No. 6025340th Globe Road, 8th Floor
      CITY: Arlington
      STATE: Virginia
      COUNTRY: USA
      COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
      CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/776.251
      FILING DATE: 31-JAN-1997
      PRIOR APPLICATION DATA:
      APPLICATION NUMBER: PCT/GB95/01782
      FILING DATE: 27-JUL-1995
      PRIOR APPLICATION DATA:
      APPLICATION NUMBER: GB 9415167.7
      FILING DATE: 27-JUL-1994
      ATTORNEY/AGENT INFORMATION:
      NAME: Arthur R. Crawford
      REGISTRATION NUMBER: 25,327
      REFERENCE/DOCKET NUMBER: 620-20
      INFORMATION FOR SEQ ID NO: 10:
      SEQUENCE CHARACTERISTICS:
      LENGTH: 816 base pairs
      TYPE: nucleic acid
      STRANDEDNESS: single
      TOPOLOGY: linear
      MOLECULE TYPE: CDNA
      US-08-776-251-10

      Query Match      73.3%; Score 15.4; DB 3; Length 816;
      Best Local Similarity 94.1%; Pred. No. 31;
      Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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OY 4 TCGAACGTTGAGATGA 20  
| | | | | | | | | |  
DB 630 TCGAACGTTGAGACGA 614

RESULT 5  
US-09-555-000-1/C  
; Sequence 1, Application US/09555000  
; Patent No. 6489108  
; GENERAL INFORMATION:  
; APPLICANT: Genecor International, Inc.  
; TITLE OF INVENTION: Proteases from Gram positive organisms  
; FILE REFERENCE: GC390-PCT  
; CURRENT APPLICATION NUMBER: US/09/555,000  
; CURRENT FILING DATE: 2000-05-23  
; PRIOR APPLICATION NUMBER: PCT/US98/26971  
; PRIOR FILING DATE: 1998-12-17  
; NUMBER OF SEQ ID NOS: 2  
; SOFTWARE: FASTSEQ for Windows Version 3.0  
; SEQ ID NO 1  
; LENGTH: 1900  
; TYPE: DNA  
; ORGANISM: Bacillus subtilis  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: (134)...(1774)  
US-09-555-000-1

Query Match 73.3% Score 15.4; DB 4; Length 1900;  
Best Local Similarity 94.1%; Pred. No. 34;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 CGAACGTTGAGATGAT 21  
| | | | | | | | | |  
DB 1119 CGAACGTTGAGATGAT 1103

RESULT 6  
US-09-134-000C-2551  
; Sequence 2551, Application US/09134000C  
; Patent No. 6617156  
; GENERAL INFORMATION:  
; APPLICANT: Lynn Doucette-Stamm et al  
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO  
; FILE REFERENCE: 032796-032  
; CURRENT APPLICATION NUMBER: US/09/134,000C  
; CURRENT FILING DATE: 1998-08-13  
; PRIOR APPLICATION NUMBER: US 60/055,778  
; PRIOR FILING DATE: 1997-08-15  
; NUMBER OF SEQ ID NOS: 6812  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 2551  
; LENGTH: 342  
; TYPE: DNA  
; ORGANISM: Enterococcus faecalis  
US-09-134-000C-2551

Query Match 72.4% Score 15.2; DB 4; Length 342;  
Best Local Similarity 85.0%; Pred. No. 36;  
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 TCGTGAACGTTGAGATGA 20  
| | | | | | | | | |  
DB 31 TCGTGAACGTTGAGATGA 50

RESULT 7  
US-09-134-001C-1108/C  
; Sequence 1108, Application US/09134001C  
; Patent No. 6380370  
; GENERAL INFORMATION:  
; APPLICANT: Raz, Eyal

APPLICANT: Lynn Doucette-Stamm et al  
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO STAPHYLOCOCCUS  
; FILE REFERENCE: GTC-007  
; CURRENT APPLICATION NUMBER: US/09/134,001C  
; CURRENT FILING DATE: 1998-08-13  
; PRIOR APPLICATION NUMBER: US 60/064,964  
; PRIOR FILING DATE: 1997-11-08  
; PRIOR APPLICATION NUMBER: US 60/055,779  
; PRIOR FILING DATE: 1997-08-14  
; NUMBER OF SEQ ID NOS: 5674  
; SEQ ID NO 1108  
; LENGTH: 4590  
; TYPE: DNA  
; ORGANISM: Staphylococcus epidermidis  
US-09-134-001C-1108

Query Match 72.4% Score 15.2; DB 4; Length 4590;  
Best Local Similarity 85.0%; Pred. No. 50;  
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 2 CGTGAACGTTGAGATGAT 21  
| | | | | | | | | |  
DB 3597 CGTGAACGTTGAGATGAT 3578

RESULT 8  
US-10-204-708-4  
; Sequence 4, Application US/10204708  
; Patent No. 6677731  
; GENERAL INFORMATION:  
; APPLICANT: OLEK, Alexander  
; APPLICANT: PIEPERROCK, Christian  
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication  
; FILE REFERENCE: 5013.1012  
; CURRENT APPLICATION NUMBER: US/10/204,708  
; CURRENT FILING DATE: 2003-05-06  
; PRIOR APPLICATION NUMBER: PCT/EP01/03971  
; PRIOR FILING DATE: 2001-04-06  
; PRIOR APPLICATION NUMBER: DE 10019058.8  
; PRIOR FILING DATE: 2000-04-06  
; PRIOR APPLICATION NUMBER: DE 10019173.8  
; PRIOR FILING DATE: 2000-04-07  
; PRIOR APPLICATION NUMBER: DE 10032529.7  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: DE 10043826.1  
; PRIOR FILING DATE: 2000-09-01  
; NUMBER OF SEQ ID NOS: 98  
; SEQ ID NO 4  
; LENGTH: 10619  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)  
US-10-204-708-4

Query Match 72.4% Score 15.2; DB 4; Length 10619;  
Best Local Similarity 85.0%; Pred. No. 55;  
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 2 CGTGAACGTTGAGATGAT 21  
| | | | | | | | | |  
DB 9684 CGTGAACGTTGAGATGAT 9703

RESULT 9  
US-09-235-742-19  
; Sequence 19, Application US/09235742  
; Patent No. 6498148  
; GENERAL INFORMATION:  
; APPLICANT: Raz, Eyal

```

; TITLE OF INVENTION: Immunization-Free Methods for Treating
; TITLE OF INVENTION: Antigen-Stimulated Inflammation in a Mammalian Host and
; TITLE OF INVENTION: Shifting the Host's Antigen Immune Responsiveness to a TH1
; FILE REFERENCE: 6510-170CON4
; CURRENT APPLICATION NUMBER: US/09/235,742
; CURRENT FILING DATE: 1999-01-21
; EARLIER APPLICATION NUMBER: 08/927,120
; EARLIER FILING DATE: 1997-09-05
; EARLIER APPLICATION NUMBER: 08/593,554
; EARLIER FILING DATE: 1996-01-30
; EARLIER APPLICATION NUMBER: 08/725,968
; EARLIER FILING DATE: 1996-10-04
; EARLIER APPLICATION NUMBER: 60/028,118
; EARLIER FILING DATE: 1996-10-11
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 19
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Recombinant or Synthetic Sequence
US-09-235-742-19

Query Match          71.4%; Score 15; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      6 GAACGTTGAGATGA 20
      |||||
Db      8 GAACGTTGAGATGA 22

RESULT 10
US-09-347-343-32
; Sequence 32, Application US/09347343A
; Patent No. 6514948
; GENERAL INFORMATION:
; APPLICANT: RAZ, Eyal R.
; APPLICANT: KOBAYASHI, HIROKO
; TITLE OF INVENTION: METHOD FOR ENHANCING AN IMMUNE RESPONSE
; FILE REFERENCE: 30448.64US01
; CURRENT APPLICATION NUMBER: US/09/347,343A
; CURRENT FILING DATE: 1999-07-02
; NUMBER OF SEQ ID NOS: 40
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 32
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Synthetic oligonucleotide
US-09-347-343-32

Query Match          71.4%; Score 15; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      6 GAACGTTGAGATGA 20
      |||||
Db      8 GAACGTTGAGATGA 22

RESULT 11
US-09-820-484-1
; Sequence 1, Application US/09820484
; Patent No. 6534062
; GENERAL INFORMATION:
; APPLICANT: RAZ, Eyal
; APPLICANT: CHO, HEARN JAY
; APPLICANT: RICHMAN, DOUGLAS
; APPLICANT: HORNER, ANTHONY A.
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
; TITLE OF INVENTION: Lymphocyte Response in vivo.
```

```

; FILE REFERENCE: 06510-188US1
; CURRENT APPLICATION NUMBER: US/09/820,484
; CURRENT FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US 60/192,537
; PRIOR FILING DATE: 2000-03-28
; PRIOR APPLICATION NUMBER: US 60/203,567
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 60/215,895
; PRIOR FILING DATE: 2000-07-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Disulfide-linked phosphorothioate ISS-ODN
; NAME/KEY: modified base
; LOCATION: (1)...(1)
; OTHER INFORMATION: disulfide thymine
US-09-820-484-1

Query Match          71.4%; Score 15; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      6 GAACGTTGAGATGA 20
      |||||
Db      8 GAACGTTGAGATGA 22

RESULT 12
US-09-820-484-3
; Sequence 3, Application US/09820484
; Patent No. 6534062
; GENERAL INFORMATION:
; APPLICANT: RAZ, Eyal
; APPLICANT: CHO, HEARN JAY
; APPLICANT: RICHMAN, DOUGLAS
; APPLICANT: HORNER, ANTHONY A.
; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
; TITLE OF INVENTION: Lymphocyte Response in vivo.
; FILE REFERENCE: 06510-188US1
; CURRENT APPLICATION NUMBER: US/09/820,484
; CURRENT FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US 60/192,537
; PRIOR FILING DATE: 2000-03-28
; PRIOR APPLICATION NUMBER: US 60/203,567
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 60/215,895
; PRIOR FILING DATE: 2000-07-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: phosphorothioate ISS-ODN
US-09-820-484-3

Query Match          71.4%; Score 15; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      6 GAACGTTGAGATGA 20
      |||||
Db      8 GAACGTTGAGATGA 22

RESULT 13
US-09-774-403A-1
; Sequence 1, Application US/09774403A
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Patent No. 6552006  
; GENERAL INFORMATION:  
; APPLICANT: Eyal Raz  
; APPLICANT: Richard Kornbluth  
; APPLICANT: Antonio Catanzaro  
; APPLICANT: Tomoko Hayashi  
; APPLICANT: Dennis Carson  
; TITLE OF INVENTION: Immunomodulatory polynucleotides in  
; treatment of infection by an intracellular pathogen  
; FILE REFERENCE: UCA1166  
; CURRENT APPLICATION NUMBER: US/09/774,403A  
; CURRENT FILING DATE: 2002-04-15  
; PRIOR APPLICATION NUMBER: 60/179,353  
; PRIOR FILING DATE: 2000-01-31  
; NUMBER OF SEQ ID NOS: 7  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Immunomodulatory sequence  
US-09-774-403A-1

Query Match 71.4%; Score 15; DB 4; Length 22;  
Best Local Similarity 100.0%; Pred. No. 32;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 GAACGTTGAGATGA 20  
|||  
DB 8 GAACGTTGAGATGA 22

RESULT 14  
US-09-296-477-1  
; Sequence 1, Application US/09296477A  
; Patent No. 6589940  
; GENERAL INFORMATION:  
; APPLICANT: RAZ, E.  
; APPLICANT: SCHWARTZ, D.  
; APPLICANT: ROMAN, M.  
; APPLICANT: DINA, D.  
; TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,  
; COMPOSITIONS THEREOF AND METHODS OF USE  
; TITLE OF INVENTION: THEREOF  
; FILE REFERENCE: 3778200420  
; CURRENT APPLICATION NUMBER: US/09/296,477A  
; CURRENT FILING DATE: 1999-04-22  
; EARLIER APPLICATION NUMBER: 09/092,329  
; EARLIER FILING DATE: 1998-06-05  
; EARLIER APPLICATION NUMBER: 60/048,793  
; EARLIER FILING DATE: 1997-06-06  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 1  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-09-296-477-1

Query Match 71.4%; Score 15; DB 4; Length 22;  
Best Local Similarity 100.0%; Pred. No. 32;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 GAACGTTGAGATGA 20  
|||  
DB 8 GAACGTTGAGATGA 22

RESULT 15  
US-09-296-477-2

Sequence 2, Application US/09296477A  
; Patent No. 6589940  
; GENERAL INFORMATION:  
; APPLICANT: RAZ, E.  
; APPLICANT: SCHWARTZ, D.  
; APPLICANT: ROMAN, M.  
; APPLICANT: DINA, D.  
; TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,  
; COMPOSITIONS THEREOF AND METHODS OF USE  
; TITLE OF INVENTION: THEREOF  
; FILE REFERENCE: 3778200420  
; CURRENT APPLICATION NUMBER: US/09/296,477A  
; CURRENT FILING DATE: 1999-04-22  
; EARLIER APPLICATION NUMBER: 09/092,329  
; EARLIER FILING DATE: 1998-06-05  
; EARLIER APPLICATION NUMBER: 60/048,793  
; EARLIER FILING DATE: 1997-06-06  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 2  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-09-296-477-2

Query Match 71.4%; Score 15; DB 4; Length 22;  
Best Local Similarity 100.0%; Pred. No. 32;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 GAACGTTGAGATGA 20  
|||  
DB 8 GAACGTTGAGATGA 22

RESULT 16  
US-09-296-477-5/c  
; Sequence 5, Application US/09296477A  
; Patent No. 6589940  
; GENERAL INFORMATION:  
; APPLICANT: RAZ, E.  
; APPLICANT: SCHWARTZ, D.  
; APPLICANT: ROMAN, M.  
; APPLICANT: DINA, D.  
; TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,  
; COMPOSITIONS THEREOF AND METHODS OF USE  
; TITLE OF INVENTION: THEREOF  
; FILE REFERENCE: 3778200420  
; CURRENT APPLICATION NUMBER: US/09/296,477A  
; CURRENT FILING DATE: 1999-04-22  
; EARLIER APPLICATION NUMBER: 09/092,329  
; EARLIER FILING DATE: 1998-06-05  
; EARLIER APPLICATION NUMBER: 60/048,793  
; EARLIER FILING DATE: 1997-06-06  
; NUMBER OF SEQ ID NOS: 21  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 5  
; LENGTH: 22  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Synthetic construct  
US-09-296-477-5

Query Match 71.4%; Score 15; DB 4; Length 22;  
Best Local Similarity 100.0%; Pred. No. 32;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 GAACGTTGAGATGA 20  
|||  
DB 15 GAACGTTGAGATGA 1

```
RESULT 17
US-09-308-036A-1
; Sequence 1, Application US/09308036A
; Patent No. 6610661
; GENERAL INFORMATION:
; APPLICANT: Carson, Dennis A.
; APPLICANT: Raz, Eyal
; APPLICANT: Roman, Mark
; TITLE OF INVENTION: Immunostimulatory
; TITLE OF INVENTION: Polynucleotide/Immunomodulatory Molecule Conjugates
; FILE REFERENCE: 6510-172CIP
; CURRENT APPLICATION NUMBER: US/09/308,036A
; CURRENT FILING DATE: 2000-02-16
; PRIOR APPLICATION NUMBER: PCT/US97/19004
; PRIOR FILING DATE: 1997-10-09
; PRIOR APPLICATION NUMBER: 60/028,118
; PRIOR FILING DATE: 1996-10-11
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO: 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: DY1018 polynucleotide
US-09-308-036A-1

Query Match      71.4%; Score 15; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      6 GAACGTTGAGATGA 20
DB      8 GAACGTTGAGATGA 22

RESULT 18
US-09-791-500-1
; Sequence 1, Application US/09791500
; Patent No. 6613751
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; APPLICANT: Rachmilewitz, Daniel
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel
; TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT APPLICATION NUMBER: US/09/791,500
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO: 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic polynucleotide sequence
US-09-791-500-1

Query Match      71.4%; Score 15; DB 4; Length 22;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      6 GAACGTTGAGATGA 20
DB      8 GAACGTTGAGATGA 22

RESULT 19
US-09-107-532A-1566
; Sequence 1566, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
```

```
APPLICANT: Lynn A Doucette-Stamm and David Bush
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
NUMBER OF SEQUENCES: 7310
CORRESPONDENCE ADDRESS:
ADDRESSER: GENOME THERAPEUTICS CORPORATION
STREET: 100 Beaver Street
CITY: Waltham
STATE: Massachusetts
COUNTRY: USA
ZIP: 02354
COMPUTER READABLE FORM:
MEDIUM TYPE: CD-ROM ISO9660
COMPUTER: PC
OPERATING SYSTEM: <Unknown>
SOFTWARE: ASCII
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/107,532A
FILING DATE: 30-Jun-1998
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/085,598
FILING DATE: 14 May 1998
APPLICATION NUMBER: 60/051571
FILING DATE: July 2, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Ariniello, Pamela Deneke
REGISTRATION NUMBER: 40,489
REFERENCE/DOCKET NUMBER: GTC-012
TELECOMMUNICATION INFORMATION:
TELEPHONE: (781)893-5007
TELEFAX: (781)893-8277
INFORMATION FOR SEQ ID NO: 1566:
SEQUENCE CHARACTERISTICS:
LENGTH: 813 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: circular
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Enterococcus faecium
FEATURE:
NAME/KEY: misc feature
LOCATION: (B) LOCATION 1...813
SEQUENCE DESCRIPTION: SEQ ID NO: 1566:
US-09-107-532A-1566

Query Match      70.5%; Score 14.8; DB 4; Length 813;
Best Local Similarity 88.9%; Pred. No. 66;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 TCGTCGACGTTGAGAT 18
DB      239 TTGTCGACGTTGGAAT 256

RESULT 20
US-09-374-174B-1
; Sequence 1, Application US/09374174B
; Patent No. 6554985
; GENERAL INFORMATION:
; APPLICANT: Ruiz-Martinez, Maria C.
; APPLICANT: Berka, Jan
; APPLICANT: Simpson, John W.
; TITLE OF INVENTION: Methods and Formulations for the Separation of
; TITLE OF INVENTION: Biological Macromolecules
; FILE REFERENCE: Cura-31: Megabace (15966-531)
; CURRENT APPLICATION NUMBER: US/09/374,174B
; CURRENT FILING DATE: 1999-08-13
; PRIOR APPLICATION NUMBER: USN 60/107,798
; PRIOR FILING DATE: 1998-11-10
; NUMBER OF SEQ ID NOS: 5
```



```

; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 1069
; TYPE: DNA
; ORGANISM: Bacteriophage M13mp18
; FEATURE:
; OTHER INFORMATION: Wherein an "n" between residues 811 to 1069 may be
; OTHER INFORMATION: G, A, T or C
US-09-374-174B-1

```

Query Match	70.5%	Score 14.8;	DB 4;	Length 1069;
Best Local Similarity	88.9%;	Pred. No. 68;		
Matches 16;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;

Qy	2	CGTCGAACGTTGAGATG	19
Db	671	CGTCGAACGTCGAGAAG	688

**RESULT 21**  
US-09-252-991A-2256  
Section 225c Analysis re/00353001A

Sequence 2256, Application US/09252991A  
Patent No. 6551795  
GENERAL INFORMATION:

: APPLICANT: Marc J. Rubenfield et al.  
 : TITLE OF INVENTION: NUCLEIC ACID SEQUENCES RELATING TO PSEUDOMONAS  
 : TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS

FILE REFERENCE: 107196.136  
CURRENT APPLICATION NUMBER: US/09/252, 991A  
CURRENT FILING DATE: 1999-02-18

;  
 ; PRIOR APPLICATION NUMBER: US 60/074,788  
 ;  
 ; PRIOR FILING DATE: 1998-02-18  
 ;  
 ; PRIOR APPLICATION NUMBER: US 60/094,190  
 ;

; PRIOR FILING DATE: 1998-07-27  
 ; NUMBER OF SEQ ID NOS: 33142  
 ; SEQ ID NO 2256

```

; SEQ ID NO 2236
; LENGTH: 1497
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa

```

```

? ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-2256

Query Match      70.5% Score 14.8 DB 4 Length 1407

```

Query Match	70.5%	Score 14.8	DB 4	Length 1487
Best Local Similarity	88.9%	Pred. No. 71		
Matches 16; Conservative	0	Mismatches 2	Indels 0	Gaps 0

```

QY      1 TCGTCGAACGTTCCGAGAT 18
          |||||
Db      1218 TCGGCGAACGATCGAGAT 1235

```

RESULT 22  
US-09-252-991A-2425/c  
Compendio 2425 and station HC/000000001A

```

; Sequence 2425, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; ADDITIONAL NOTES:

```

; APPLICANT: Marc J. Rubenfield et al.  
 ; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS  
 ; TITLE OF INVENTION: AERUCINOSA FOR DIAGNOSTICS AND THERAPEUTICS

FILE REFERENCE: 107196.136  
CURRENT APPLICATION NUMBER: US/09/252,991A  
CURRENT FILING DATE: 1999-02-18

; PRIOR APPLICATION NUMBER: US 60/074,788  
 ; PRIOR FILING DATE: 1998-02-18  
 ; PRIOR APPLICATION NUMBER: US 60/094,190

; PRIOR FILING DATE: 1998-07-27  
 ; NUMBER OF SEQ ID NOS: 33142  
 ; SEQ ID NO 2425

; ORG: AD NO 2123  
 ; LENGTH: 1950  
 ; TYPE: DNA  
 ; ORGANISM: *Pseudomonas aeruginosa*

S-09-252-991A-2425

Query Match	70.5%;	Score 14.8;	DB 4;	Length 1950;
-------------	--------	-------------	-------	--------------

Best Local Similarity 88.9%; Pred.No. 74;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

OY      1 TCCTGACGTTGAGAT 18
          ||| ||||| |||||
Db      847 TCGCGACGATCGAGAT 830

```

RESULT 23  
US-08-446-794A-5  
Sequence 5 Amplification ITS/08446794A

; Sequence 5, Application US/08446794A  
 ; Patent No. 5747327  
 ; GENERAL INFORMATION:  
 ;  
 ; ADDITIONAL INFORMATION:

APPLICANT: UEKI, JUN  
APPLICANT: MORIOKA, SHINJI  
TITLE OF INVENTION: PHOSPHOLIPASE D GENE ORIGINATED FROM

```

;
; TITLE OF INVENTION: PLANT
;
; NUMBER OF SEQUENCES: 7
;
; CORRESPONDENCE ADDRESS:
;

```

ADDRESSEE: BIRCH, ST  
STREET: P.O. BOX 74  
CITY: FALLS CHURCH

STATE: VA  
COUNTRY: USA  
ZIP: 22040-0747

```

;          DATE: 22070 0787
;
;      COMPUTER READABLE FORM
;      MEDIUM TYPE: Floppy
;      COMPUTER: IBM PC compatible

```

```

;
; COMPUTER: IBM PC Compatible
;
; OPERATING SYSTEM: PC-DOS/MS-DOS
;
; SOFTWARE: PatentIn Release #1.0, Version #1.30
;
CURRENT APPLICATION DATA:
```

CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/446,794A  
 FILING DATE:  
 CLASSIFICATION: 536

CLASSIFICATION: 536  
ATTORNEY/AGENT INFORMATION:  
NAME: MURPHY JR, GERALD M  
REGISTRATION NUMBER: 38 8

REGISTRATION NUMBER: 28,977  
REFERENCE/DOCKET NUMBER: 0760-02031  
TELECOMMUNICATION INFORMATION:  
SERIAL NUMBER: 100 00 0000

TELEPHONE: 703-205-8000  
TELEFAX: 703-205-8050  
INFORMATION FOR SEQ ID NO: 5:

```

; SEQUENCE CHARACTERISTICS:
; LENGTH: 2799 base pairs
; TYPE: nucleic acid
;

```

```

STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)

```

```

; NOBLOCED FILE: ENR (GENERAL)
;
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1876..1983

```

```

/ LOCATION: 1878.1363
/
/ FEATURE:
/ NAME/KEY: CDS
/ LOCATION: 2524.2799
/

```

```

;      LOCATION: 2524..2/99
US-08-446-794A-5
Query Match      70 5%:

```

Query Match	70.5%	Score	14.8	DB	1	Length	2759
Best Local Similarity	88.9%	Pred. No.	77				
Matches	16	Conservative	0	Mismatches	2	Indels	0
						Gaps	0

QY 3 GTGGAACGTTGAGATGA 20  
||| ||| ||| ||| ||| |||  
Db 751 GTCAACGTTGATATGA 768

RESULT 24  
US-08-750-007-4  
Sequence A Amplification US/08750007

; Sequence 4, Application US/08750007  
 ; Patent No. 5801016  
 ; GENERAL INFORMATION:  
 ;  
 ADDICANT, MODICA CHINTI

; APPLICANT: MORIOKA, SHINOBU  
 ; APPLICANT: UEKI, JUN  
 ; TITLE OF INVENTION: DNA FRAGMENT, RECOMBINANT VECTOR

TITLE OF INVENTION: CONTAINING THE SAME AND METHOD FOR EXPRESSING FOREIGN  
TITLE OF INVENTION: GENES USING THE SAME  
NUMBER OF SEQUENCES: 19  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: BIRCH, STEWART, KOLASCH AND BIRCH  
STREET: PO BOX 747  
CITY: FALLS CHURCH  
STATE: VA  
COUNTRY: USA  
ZIP: 22040-0747  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/750,007  
FILING DATE:  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: MURPHY JR, GERALD M  
REGISTRATION NUMBER: 28,977  
REFERENCE/DOCKET NUMBER: 0760-221P  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (703) 205-8000  
TELEFAX: (703) 205-8050  
INFORMATION FOR SEQ ID NO: 4:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 2799 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1876..1983  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 2524..2799  
US-08-750-007-4

Query Match 70.5%; Score 14.8; DB 1; Length 2799;  
Best Local Similarity 88.9%; Pred. No. 77;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 GTGGAACGTTGAGATGA 20  
||| ||||| |||||  
DB 751 GTCAACGTTGATATGA 768

RESULT 25  
US-09-621-976-12664  
Sequence 12664, Application US/09621976  
Patent No. 6639063  
GENERAL INFORMATION:  
APPLICANT: Dumas Milne Edwards, J.B.  
APPLICANT: Jobert, S.  
APPLICANT: Giordano, J.Y.  
TITLE OF INVENTION: ESTs and Encoded Human Proteins.  
FILE REFERENCE: GENSET.054PR2  
CURRENT APPLICATION NUMBER: US/09/621,976  
CURRENT FILING DATE: 2000-07-21  
NUMBER OF SEQ ID NOS: 19335  
SOFTWARE: Patent.pm  
SEQ ID NO 12664  
LENGTH: 424  
TYPE: DNA  
ORGANISM: Homo sapiens  
US-09-621-976-12664

Query Match 69.5%; Score 14.6; DB 4; Length 424;  
Best Local Similarity 81.0%; Pred. No. 78;  
Matches 17; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TCGTGAACGTTGAGATGAT 21  
||| ||||| ||||| |||||  
DB 115 TCGTGAACGTTGAAATGAT 135

RESULT 26  
US-09-221-017B-893  
Sequence 893, Application US/09221017B  
Patent No. 6444799  
GENERAL INFORMATION:  
APPLICANT: Ross, Bruce C.  
TITLE OF INVENTION: P. GINGIVALIS NUCLEOTIDES AND USES THEREOF  
NUMBER OF SEQUENCES: 1120  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: MORRISON & FOERSTER  
STREET: 755 PAGE MILL ROAD  
CITY: Palo Alto  
STATE: CA  
COUNTRY: USA  
ZIP: 94304-1018  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: Windows  
SOFTWARE: FastSeq for Windows Version 2.0b  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/221,017B  
FILING DATE: 23-DEC-1998  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PP1182  
FILING DATE: 31-DEC-1997  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PP1546  
FILING DATE: 30-JAN-1998  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PP2311  
FILING DATE: 09-APR-1998  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PCT/AU98/01023  
FILING DATE: 10-DEC-1998  
ATTORNEY/AGENT INFORMATION:  
NAME: Monroy, Gladys H  
REGISTRATION NUMBER: 32,430  
REFERENCE/DOCKET NUMBER: 27340-20021.00  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 650-813-5600  
TELEFAX: 650-494-0792  
TELEX: 706141  
INFORMATION FOR SEQ ID NO: 893:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3468 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: circular  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: UNKNOWN  
ORIGINAL SOURCE:  
ORGANISM: PORPHYROMONAS GINGIVALIS  
FEATURE:  
NAME/KEY: misc feature  
LOCATION: 1...3468  
US-09-221-017B-893

Query Match 69.5%; Score 14.6; DB 4; Length 3468;  
Best Local Similarity 81.0%; Pred. No. 1e+02;  
Matches 17; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TCGTGAACGTTGAGATGAT 21  
||| ||||| ||||| |||||  
DB 1936 TCGTGTGCTTTCGAGTGAT 1956

RESULT 27  
US-08-399-412A-58  
Sequence 58, Application US/08399412A  
Patent No. 5622828  
GENERAL INFORMATION:  
APPLICANT: Parma, David  
TITLE OF INVENTION: Gold, Larry  
TITLE OF INVENTION: High-Affinity Oligonucleotide  
TITLE OF INVENTION: Ligands To Secretory Phospholipase  
TITLE OF INVENTION: A2 (sPLA2)  
NUMBER OF SEQUENCES: 122  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Swanson & Bratschun, L.L.C.  
STREET: 8400 E. Prentice Avenue, Suite 200  
CITY: Englewood  
STATE: Colorado  
COUNTRY: USA  
ZIP: 80111  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB storage  
COMPUTER: IBM compatible  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WordPerfect 5.1  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/399,412A  
FILING DATE: 6-MARCH-1995  
CLASSIFICATION: 536  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/714,131  
FILING DATE: 10-JUNE-1991  
CLASSIFICATION: 536  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/536,428  
FILING DATE: 11-JUNE-1990  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/964,624  
FILING DATE: 21-OCTOBER-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: Julie L. Bernard  
REGISTRATION NUMBER: 36,450  
REFERENCE/DOCKET NUMBER: NEX27  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (303) 793-3333  
TELEFAX: (303) 793-3433  
INFORMATION FOR SEQ ID NO: 58:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 77 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
US-08-399-412A-58

Query Match 68.6%; Score 14.4; DB 1; Length 77;  
Best Local Similarity 75.0%; Pred. No. 80;  
Matches 12; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 5 CGACGCTTGAGATGA 20  
|||:|||||:|  
Db 48 CGACGCTTGAGATGA 63

RESULT 28  
US-08-447-169A-36  
Sequence 36, Application US/08447169A  
Patent No. 581533  
GENERAL INFORMATION:  
APPLICANT: JANUIC, N. and GOLD, L.  
TITLE OF INVENTION: HIGH-AFFINITY OLIGONUCLEOTIDE  
TITLE OF INVENTION: LIGANDS TO VASCULAR ENDOTHELIAL  
TITLE OF INVENTION: GROWTH FACTOR (VEGF)  
NUMBER OF SEQUENCES: 242

CORRESPONDENCE ADDRESS:  
ADDRESSEE: Swanson & Bratschun, L.L.C.  
STREET: 8400 E. Prentice Place, Suite 200  
CITY: Englewood  
STATE: Colorado  
COUNTRY: USA  
ZIP: 80111  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.50 inch, 1.44 MB storage  
COMPUTER: IBM compatible  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WordPerfect 5.1  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/447,169A  
FILING DATE: 19-MAY-1995  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/233,012  
FILING DATE: 25-APRIL-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/205,515  
FILING DATE: 03-MARCH-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/964,624  
FILING DATE: 21-OCTOBER-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/714,131  
FILING DATE: 10-JUNE-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/536,428  
FILING DATE: 11-JUNE-1990  
ATTORNEY/AGENT INFORMATION:  
NAME: Barry J. Swanson  
REGISTRATION NUMBER: 33,215  
REFERENCE/DOCKET NUMBER: NEX14  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (303) 793-3333  
TELEFAX: (303) 793-3433  
INFORMATION FOR SEQ ID NO: 36:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 77 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
US-08-447-169A-36

Query Match 67.6%; Score 14.2; DB 1; Length 77;  
Best Local Similarity 63.2%; Pred. No. 1e+02;  
Matches 12; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 2 CGTCGACGCTTGAGATGA 20  
||:|||||:|  
Db 45 CGUAGAGCGGUCGACAUCA 63

RESULT 29  
US-08-233-012C-36  
Sequence 36, Application US/08233012C  
Patent No. 5849479  
GENERAL INFORMATION:  
APPLICANT: JANUIC, N. and GOLD, L.  
TITLE OF INVENTION: HIGH-AFFINITY OLIGONUCLEOTIDE  
TITLE OF INVENTION: LIGANDS TO VASCULAR  
TITLE OF INVENTION: ENDOTHELIAL GROWTH FACTOR  
NUMBER OF SEQUENCES: 146  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Swanson & Bratschun, L.L.C.  
STREET: 8400 E. Prentice Place, Suite 200  
CITY: Englewood  
STATE: Colorado  
COUNTRY: USA  
ZIP: 80111

```
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44 MG
MEDIUM TYPE: storage
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/233,012C
FILING DATE: 25-APRIL-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/714,131
FILING DATE: 10-JUNE-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/536,428
FILING DATE: 11-JUNE-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/964,624
FILING DATE: 21-OCTOBER-1992
ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER: NEX14
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 77 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-233-012C-36

Query Match          67.6%; Score 14.2; DB 2; Length 77;
Best Local Similarity 63.2%; Pred. No. 1e+02;
Matches 12; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY      2 CGTCGACGTCGAGATGA 20
DB      45 CGUAGAGCGUCCGACAUGA 63

RESULT 30
US-09-848-295-1
; Sequence 1, Application US/09848295
; Patent No. 6623941
; GENERAL INFORMATION:
; APPLICANT: Baker, Kevin
; APPLICANT: Ruben, Steven M.
; TITLE OF INVENTION: Human Tumor Necrosis Factor TR20 and Methods Based
; TITLE OF INVENTION: Thereon
; FILE REFERENCE: PF527
; CURRENT APPLICATION NUMBER: US/09/848,295
; CURRENT FILING DATE: 2001-05-04
; PRIOR APPLICATION NUMBER: 60/202,193
; PRIOR FILING DATE: 2000-05-05
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 1535
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (154)..(579)
US-09-848-295-1

Query Match          67.6%; Score 14.2; DB 4; Length 1535;
Best Local Similarity 84.2%; Pred. No. 1.5e+02;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY      3 GTCGACGTCGAGATGAT 21
```

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DB      1467 GTCGAAGTTCGAGACCAT 1485

RESULT 31
US-08-956-171E-203
; Sequence 203, Application US/08956171E
; Patent No. 6593114
; GENERAL INFORMATION:
; APPLICANT: Charles Kunesh
;           Gail H. Choi
;           Patrick S. Dillon
;           Craig A. Rosen
;           Steven C. Barash
;           Michael R. Fannon
; TITLE OF INVENTION: Staphylococcus aureus Polynucleotides and Sequences
; NUMBER OF SEQUENCES: 5256
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20850
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44MB storage
COMPUTER: HP Vectra 486/33
OPERATING SYSTEM: MSDOS version 6.2
SOFTWARE: ASCII Text
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/956,171E
FILING DATE: 20-OCT-1997
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/009,861
FILING DATE: January 5, 1996
APPLICATION NUMBER: 08/781,986
FILING DATE: January 3, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Mark J. Hyman
REGISTRATION NUMBER: 46,789
REFERENCE/DOCKET NUMBER: PB248P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (240) 314-1224
TELEFAX: (301) 309-8439
INFORMATION FOR SEQ ID NO: 203:
SEQUENCE CHARACTERISTICS:
LENGTH: 4715 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 203:
US-08-956-171E-203

Query Match          67.6%; Score 14.2; DB 4; Length 4715;
Best Local Similarity 84.2%; Pred. No. 1.8e+02;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY      2 CGTCGACGTCGAGATGA 20
DB      1285 CGTTAAAGTTCGAGATGA 1303

RESULT 32
US-09-173-914-6/C
; Sequence 6, Application US/09173914
; Patent No. 6171857
; GENERAL INFORMATION:
; APPLICANT: Hendrickson, Eric
; TITLE OF INVENTION: A No. 6171857e1 Leucine zipper, KARP-1 and
; TITLE OF INVENTION: Methods of Regulating DNA Dependent Protein Kinase Activity
; FILE REFERENCE: B0877/7017/HK
; CURRENT APPLICATION NUMBER: US/09/173,914
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; CURRENT FILING DATE: 1998-10-16
; EARLIER APPLICATION NUMBER: 60/064,557
; EARLIER FILING DATE: 1997-10-17
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 6
; LENGTH: 14636
; TYPE: DNA
; ORGANISM: Homo Sapiens
; US-09-173-914-6

Query Match
Best Local Similarity 67.6%; Score 14.2; DB 3; Length 14636;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 CGTCGAACGTTGAGATGA 20
Db 1050 CTTCAACGTTGAGATGA 1032

RESULT 33
US-09-428-517-1/c
; Sequence 1, Application US/09428517
; Patent No. 6251636
; GENERAL INFORMATION:
; APPLICANT: Bellich, Mary C.
; APPLICANT: Shah, Sanjay Krishnakant
; APPLICANT: McDaniel, Robert
; APPLICANT: Tang, Li
; TITLE OF INVENTION: RECOMBINANT OLEANDOLIDE POLYKETIDE SYNTHASE
; FILE REFERENCE: 30062-20029.00
; CURRENT FILING DATE: 1999-10-28
; EARLIER APPLICATION NUMBER: 60/120,254
; EARLIER FILING DATE: 1999-02-16
; EARLIER APPLICATION NUMBER: 60/106,100
; EARLIER FILING DATE: 1998-10-29
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 50937
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Recombinant DNA
; US-09-428-517-1

Query Match
Best Local Similarity 67.6%; Score 14.2; DB 3; Length 50937;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 TCGTGAACGTTGAGATG 19
Db 5439 TCGTGAACGTTGAGATG 5421

RESULT 34
US-09-103-840A-2/c
; Sequence 2, Application US/09103840A
; Patent No. 6294328
; GENERAL INFORMATION:
; APPLICANT: FLEISCHMAN, Robert D.
; APPLICANT: WHITE, Owen R.
; APPLICANT: FRASER, Claire M.
; APPLICANT: VENTER, John C.
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM
; TITLE OF INVENTION: TUBERCULOSIS
; FILE REFERENCE: 24366-20007.00
; CURRENT FILING DATE: 1998-06-24
; EARLIER APPLICATION NUMBER: 60/048,793
; EARLIER FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
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```

; LENGTH: 4403765
; TYPE: DNA
; ORGANISM: Mycobacterium tuberculosis
; FEATURE:
; OTHER INFORMATION: CDC 1551
; OTHER INFORMATION: "n" bases at various positions throughout the sequence
; OTHER INFORMATION: represent a, t, c or g
; US-09-103-840A-2

Query Match
Best Local Similarity 67.6%; Score 14.2; DB 3; Length 4403765;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 GTCCGAACGTTGAGATGAT 21
Db 1399888 GTCCGCCGTTGAGATGAT 1399870

RESULT 35
US-09-103-840A-1/c
; Sequence 1, Application US/09103840A
; Patent No. 6294328
; GENERAL INFORMATION:
; APPLICANT: FLEISCHMAN, Robert D.
; APPLICANT: WHITE, Owen R.
; APPLICANT: FRASER, Claire M.
; APPLICANT: VENTER, John C.
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM
; TITLE OF INVENTION: TUBERCULOSIS
; FILE REFERENCE: 24366-20007.00
; CURRENT FILING DATE: 1998-06-24
; EARLIER APPLICATION NUMBER: 60/048,793
; EARLIER FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 4411529
; TYPE: DNA
; ORGANISM: Mycobacterium tuberculosis
; OTHER INFORMATION: H37Rv
; US-09-103-840A-1

Query Match
Best Local Similarity 67.6%; Score 14.2; DB 3; Length 4411529;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 GTCCGAACGTTGAGATGAT 21
Db 1400419 GTCCGCCGTTGAGATGAT 1400401

RESULT 36
US-09-296-477-15
; Sequence 15, Application US/09296477A
; Patent No. 6589940
; GENERAL INFORMATION:
; APPLICANT: RAZ, E.
; APPLICANT: SCHWARTZ, D.
; APPLICANT: ROMAN, M.
; APPLICANT: DINA, D.
; TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES,
; TITLE OF INVENTION: COMPOSITIONS THEREOF AND METHODS OF USE
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: 37782000420
; CURRENT FILING DATE: 1999-04-22
; EARLIER APPLICATION NUMBER: 09/092,329
; EARLIER FILING DATE: 1998-06-05
; EARLIER APPLICATION NUMBER: 60/048,793
; EARLIER FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 15
; LENGTH: 22
```

TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Synthetic construct  
FEATURE:  
NAME/KEY: modified base  
LOCATION: (11)-(11)  
OTHER INFORMATION: 5-bromocytosine  
US-09-296-477-15

Query Match 66.7%; Score 14; DB 4; Length 22;  
Best Local Similarity 93.3%; Pred. No. 1.1e+02;  
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 GAACGTTGAGATGA 20  
DB 8 GAAGTTGAGATGA 22

RESULT 37  
US-08-710-134-63/c  
Sequence 63, Application US/08710134  
Patent No. 5834181  
GENERAL INFORMATION:  
APPLICANT: SHUBER, ANTHONY P.  
TITLE OF INVENTION: HIGH THROUGHPUT SCREENING METHOD FOR  
TITLE OF INVENTION: SEQUENCES OR GENETIC ALTERATIONS IN NUCLEIC ACIDS  
NUMBER OF SEQUENCES: 65  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genzyme Corporation  
STREET: One Mountain Road  
CITY: Framingham  
STATE: Massachusetts  
COUNTRY: USA  
ZIP: 01701  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/710,134  
FILING DATE: 13-SEP-1996  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Dugan, Deborah A.  
REGISTRATION NUMBER: 37,315  
REFERENCE/DOCKET NUMBER: IGS-8.1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 508-872-8400  
TELEFAX: 508-872-5415  
INFORMATION FOR SEQ ID NO: 63:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 66 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: other nucleic acid  
DESCRIPTION: /desc = "Oligonucleotides"  
US-08-710-134-63

Query Match 65.7%; Score 13.8; DB 2; Length 66;  
Best Local Similarity 88.2%; Pred. No. 1.7e+02;  
Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 TCGACGTTGAGATGA 20  
DB 51 TCGATGATGAGATGA 35

RESULT 38  
US-08-485-885-63/c  
Sequence 63, Application US/08485885

Patent No. 5849483  
GENERAL INFORMATION:  
APPLICANT: SHUBER, ANTHONY P.  
TITLE OF INVENTION: HIGH THROUGHPUT SCREENING METHOD FOR  
TITLE OF INVENTION: SEQUENCES OR GENETIC ALTERATIONS IN NUCLEIC ACIDS  
NUMBER OF SEQUENCES: 65  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genzyme Corporation  
STREET: One Mountain Road  
CITY: Framingham  
STATE: Massachusetts  
COUNTRY: USA  
ZIP: 01701  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/485,885  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Dugan, Deborah A.  
REGISTRATION NUMBER: 37,315  
REFERENCE/DOCKET NUMBER: GEN4-12.1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 508-872-8400  
TELEFAX: 508-872-5415  
INFORMATION FOR SEQ ID NO: 63:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 66 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: other nucleic acid  
DESCRIPTION: /desc = "Oligonucleotides"  
US-08-485-885-63

Query Match 65.7%; Score 13.8; DB 2; Length 66;  
Best Local Similarity 88.2%; Pred. No. 1.7e+02;  
Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 TCGACGTTGAGATGA 20  
DB 51 TCGATGATGAGATGA 35

RESULT 39  
US-08-710-134-64/c  
Sequence 64, Application US/08710134  
Patent No. 5834181  
GENERAL INFORMATION:  
APPLICANT: SHUBER, ANTHONY P.  
TITLE OF INVENTION: HIGH THROUGHPUT SCREENING METHOD FOR  
TITLE OF INVENTION: SEQUENCES OR GENETIC ALTERATIONS IN NUCLEIC ACIDS  
NUMBER OF SEQUENCES: 65  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genzyme Corporation  
STREET: One Mountain Road  
CITY: Framingham  
STATE: Massachusetts  
COUNTRY: USA  
ZIP: 01701  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/710,134  
FILING DATE: 13-SEP-1996  
CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:  
 NAME: Dugan, Deborah A.  
 REGISTRATION NUMBER: 37,315  
 REFERENCE/DOCKET NUMBER: 1G5-8.1  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: 508-872-8400  
 TELEFAX: 508-872-5415  
 INFORMATION FOR SEQ ID NO: 64:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 86 base pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: other nucleic acid  
 DESCRIPTION: /desc = "Oligonucleotides"  
 US-08-710-134-64

Query Match 65.7%; Score 13.8; DB 2; Length 86;  
 Best Local Similarity 88.2%; Pred. No. 1.7e+02;  
 Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 TCGAAGTTGAGATGA 20  
 |||||  
 Db 70 TCGATCGATCGAGATGA 54

RESULT 40  
 US-08-485-885-64/c  
 Sequence 64, Application US/08485885  
 Patent No. 5849483  
 GENERAL INFORMATION:  
 APPLICANT: SHUBER, ANTHONY P.  
 TITLE OF INVENTION: HIGH THROUGHPUT SCREENING METHOD FOR  
 TITLE OF INVENTION: SEQUENCES OR GENETIC ALTERATIONS IN NUCLEIC ACIDS  
 NUMBER OF SEQUENCES: 65  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Genzyme Corporation  
 STREET: One Mountain Road  
 CITY: Framingham  
 STATE: Massachusetts  
 COUNTRY: USA  
 ZIP: 01701  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 SOFTWARE: PatentIn Release #1.0, Version #1.30  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/485,885  
 FILING DATE: 07-JUN-1995  
 CLASSIFICATION: 435  
 ATTORNEY/AGENT INFORMATION:  
 NAME: Dugan, Deborah A.  
 REGISTRATION NUMBER: 37,315  
 REFERENCE/DOCKET NUMBER: GEN4-12.1  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: 508-872-8400  
 TELEFAX: 508-872-5415  
 INFORMATION FOR SEQ ID NO: 64:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 86 base pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: other nucleic acid  
 DESCRIPTION: /desc = "Oligonucleotides"  
 US-08-485-885-64

Query Match 65.7%; Score 13.8; DB 2; Length 86;  
 Best Local Similarity 88.2%; Pred. No. 1.7e+02;  
 Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 TCGAAGTTGAGATGA 20

Db 70 TCGATCGATCGAGATGA 54  
 |||||

Search completed: April 6, 2004, 03:58:41  
 Job time: 114 secs

**This Page Blank (uspto)**



GenCore version 5.1.6  
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# OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 01:36:06 ; Search time 3183 Seconds  
(without alignments)  
197.017 Million cell updates/sec

Title: US-10-033-243-132

Perfect score: 21

Sequence: 1 tcgtcgaactcgaatgat 21

Scoring table: IDENTITY NUC  
Gapop 10.0, Gapext 1.0

Searched: 27513289 seqs, 14931090276 residues  
Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database :

EST:  
1: em\_eebba:\*  
2: em\_eebba:\*  
3: em\_eebba:\*  
4: em\_eebba:\*  
5: em\_eebba:\*  
6: em\_eebba:\*  
7: em\_eebba:\*  
8: em\_eebba:\*  
9: gb\_eeb1:\*  
10: gb\_eeb2:\*  
11: gb\_eeb3:\*  
12: gb\_eeb3:\*  
13: gb\_eeb3:\*  
14: gb\_eeb3:\*  
15: em\_eebba:\*  
16: em\_eebba:\*  
17: em\_gss\_hum:\*  
18: em\_gss\_hum:\*  
19: em\_gss\_hum:\*  
20: em\_gss\_hum:\*  
21: em\_gss\_hum:\*  
22: em\_gss\_hum:\*  
23: em\_gss\_hum:\*  
24: em\_gss\_hum:\*  
25: em\_gss\_hum:\*  
26: em\_gss\_hum:\*  
27: em\_gss\_hum:\*  
28: gb\_gss2:\*  
29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	18.4	87.6	802	29	CNS04SNV
2	17.8	84.8	302	14	CD182992
3	17.8	84.8	441	14	CD156376
4	17.8	84.8	586	14	CD124255

C	5	17.8	84.8	785	14	CF434519	CF434519	EST670864
C	6	16.8	80.0	221	14	CD572389	CD572389	PBL_20_H0
C	7	16.8	80.0	461	14	CA707068	CA707068	wk2c_PX0
C	8	16.8	80.0	587	14	CA827269	CA827269	1114012C1
C	9	16.8	80.0	595	28	BH387860	BH387860	AG-ND-129
C	10	16.8	80.0	1410	10	BE052250	BE052250	GA_Ea003
C	11	16.4	78.1	303	10	AW710594	AW710594	e4D10ne.x
C	12	16.4	78.1	424	13	BM170465	BM170465	BM170465
C	13	16.4	78.1	439	12	BM158988	BM158988	BM160020B
C	14	16.4	78.1	445	13	BM300285	BM300285	BM300285
C	15	16.4	78.1	457	12	BP018564	BP018564	BP018564
C	16	16.4	78.1	504	28	B29433	B29433	F1912TFB_I
C	17	16.4	78.1	511	12	B1514876	B1514876	BM160016B
C	18	16.4	78.1	511	12	B1514893	B1514893	BM160016B
C	19	16.4	78.1	635	13	BM18728	BM18728	BM18728
C	20	16.4	78.1	650	9	AV902415	AV902415	AV902415
C	21	16.4	78.1	662	13	BM174317	BM174317	BM174317
C	22	16.4	78.1	663	13	BM171245	BM171245	BM171245
C	23	16.4	78.1	707	13	BM297679	BM297679	BM297679
C	24	16.4	78.1	709	13	BM303599	BM303599	BM303599
C	25	16.4	78.1	726	13	BM174467	BM174467	BM174467
C	26	16.4	78.1	914	29	CNS04A00	CNS04A00	AL282105
C	27	16.2	77.1	166	14	CA196886	CA196886	SCBFD109
C	28	16.2	77.1	273	12	BG931701	BG931701	1112-118
C	29	16.2	77.1	314	14	CD190559	CD190559	MS1-0064U
C	30	16.2	77.1	315	14	CD183371	CD183371	MS1-0038U
C	31	16.2	77.1	328	12	BI075646	BI075646	IP1_23_CO
C	32	16.2	77.1	329	14	CD177754	CD177754	MS1-0011T
C	33	16.2	77.1	342	14	CD092357	CD092357	MC1-0096T
C	34	16.2	77.1	374	28	AQ840347	AQ840347	ncbX0055D
C	35	16.2	77.1	375	14	CD190376	CD190376	MS1-0064U
C	36	16.2	77.1	377	14	CD178552	CD178552	MS1-0014P
C	37	16.2	77.1	384	14	CD187006	CD187006	MS1-0056U
C	38	16.2	77.1	384	14	CD190545	CD190545	MS1-0064U
C	39	16.2	77.1	392	14	CD124011	CD124011	MS1-0084T
C	40	16.2	77.1	410	10	AW699669	AW699669	gb30d0c.Y
C	41	16.2	77.1	412	10	AW497269	AW497269	gb57c10.Y
C	42	16.2	77.1	414	14	CD096098	CD096098	ME1-0006T
C	43	16.2	77.1	428	14	BU195526	BU195526	BU195526
C	44	16.2	77.1	428	12	CF431104	CF431104	NT11_6_D1
C	45	16.2	77.1	434	14	CF431883	CF431883	NT11_11_E

## ALIGNMENTS

RESULT 1	CNS04SNV	802 bp	DNA	linear	GSS 01-SEP-2000
LOCUS	Tetradodon nigroviridis genome survey sequence T7 end of clone				
DEFINITION	00710 of library H from Tetradodon nigroviridis, genomic survey sequence.				
ACCESSION	AL305428.1	GI:8197678			
VERSION	AL305428				
KEYWORDS	GSS: genome survey sequence.				
SOURCE	Tetradodon nigroviridis				
ORGANISM	Tetradodon nigroviridis				
REFERENCE	Roest Crolius,H., Jallion,O., Dasilva,C., Bouneau,L., Fisher,C., Bernot,A., Fizames,C., Winkler,P., Brotlier,P., Quetier,F., Saurin,W. and Weissenbach,J.				
AUTHORS	Estimate of human gene number provided by genome-wide analysis using Tetradodon nigroviridis DNA sequence				
TITLE	Nat. Genet. 25 (2), 235-238 (2000)				
JOURNAL	20296633				
MEDLINE	10835645				
PUBMED					
REFERENCE	Roest Crolius,H., Jallion,O., Dasilva,C., Ozouf-Costaz,C., Fizames,C., Fischer,C., Bouneau,L., Billault,A., Quetier,F.,				
AUTHORS					

**TITLE** Saurin, W., Bernot, A. and Weissbach, J.  
**CHARACTERIZATION AND REPEAT ANALYSIS OF THE COMPACT GENOME OF THE FRESHWATER PUFFERFISH TETRAODON NIGROVIRIDIS**  
**JOURNAL** Genome Res. 10 (7), 939-949. (2000)  
**MEDLINE** 20359837  
**PUBMED** 10899143  
**REFERENCE** 3 (bases 1 to 802)  
**AUTHORS** Genoscope.  
**TITLE** Direct Submission  
**JOURNAL** Submitted (12-APR-2000) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : sequef@genoscope.cns.fr - Web : www.genoscope.cns.fr)  
 This sequence is a single read and was generated as part of a large scale clone-end sequencing project of the Tetraodon nigroviridis genome. For more information, please take a look at <http://www.genoscope.cns.fr/Tetraodon>.  
**FEATURES** Location/Qualifiers  
 source  
 1..802  
 /organism="Tetraodon nigroviridis"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:99883"  
 /clone="007010"  
 /clone\_1lb="H"  
 /note="Genoscope sequence ID : C0BH007DE05XD1-end : T7"

**ORIGIN**  
 Query Match 87.6%; Score 18.4; DB 29; Length 802;  
 Best Local Similarity 95.0%; Pred. No. 2.6e+02;  
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

**QY** 2 CGTCGACGCTGCAGATGAT 21  
 |||||  
 623 CGTCGACGCTGCAGATGAT 642

**RESULT 2**  
**LOCUS** CD182992 302 bp mRNA linear EST 14-SEP-2003  
**DEFINITION** MS1-0037T-D120-A09-U-G MS1-0037 Schistosoma mansoni cDNA clone  
**ACCESSION** CD182992  
**VERSION** CD182992.1 GI:34713214  
**KEYWORDS** EST.  
**SOURCE** Schistosoma mansoni  
**ORGANISM** Schistosoma mansoni  
 Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae; Schistosoma.  
**REFERENCE** 1 (bases 1 to 302)  
 Verjovski-Almeida, S., Demarco, R., Martins, E.A.L., Guimaraes, P.E.M., Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y., Jr., Kitajima, J.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F., Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L., Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A., Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A., Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T., Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M., Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.  
**TITLE** Transcriptome analysis of the acclimated human parasite Schistosoma mansoni  
**JOURNAL** Nat. Genet. 35 (2), 148-157 (2003)  
**MEDLINE** 22879926  
**COMMENT** Contact: Dr. Sergio Verjovski-Almeida  
 Departamento de Bioquimica  
 Instituto de Quimica - Universidade de Sao Paulo  
 Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 Sao Paulo - SP, Brasil  
 Tel: +55-11-3091-2173  
 Fax: +55-11-3091-2186  
 Email: verjov@iq.usp.br  
 This sequence was derived from the FAPESP Schistosoma mansoni EST Genome Project. All sequences in the project were assembled and annotated. This entry and all the assembled sequences can be seen in the following URL <http://bioinfo.iq.usp.br/schisto/>

**FEATURES** Location/Qualifiers  
 source  
 1..302  
 /organism="Schistosoma mansoni"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:6183"  
 /clone="MS1-0037T-D120-A09.G"  
 /sex="mixed pool"  
 /dev\_stage="schistosomulum"  
 /lab\_host="in vitro culture"  
 /clone\_1lb="MS1-0037"  
 /note="Vector: pGEM T-easy"

**ORIGIN**  
 Query Match 84.8%; Score 17.8; DB 14; Length 302;  
 Best Local Similarity 90.5%; Pred. No. 3.8e+02;  
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

**QY** 1 TCCTCGACGCTGCAGATGAT 21  
 |||||  
 19 TCCTCGACGCTGTGATGAT 39

**RESULT 3**  
**LOCUS** CD156376/c 441 bp mRNA linear EST 14-SEP-2003  
**DEFINITION** ML1-0046T-M209-C11-U-G ML1-0046 Schistosoma mansoni cDNA clone  
**ACCESSION** CD156376  
**VERSION** CD156376.1 GI:34693161  
**KEYWORDS** EST.  
**SOURCE** Schistosoma mansoni  
**ORGANISM** Schistosoma mansoni  
 Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae; Schistosoma.  
**REFERENCE** 1 (bases 1 to 441)  
 Verjovski-Almeida, S., Demarco, R., Martins, E.A.L., Guimaraes, P.E.M., Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y., Jr., Kitajima, J.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F., Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L., Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A., Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A., Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T., Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M., Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.  
**TITLE** Transcriptome analysis of the acclimated human parasite Schistosoma mansoni  
**JOURNAL** Nat. Genet. 35 (2), 148-157 (2003)  
**MEDLINE** 22879926  
**COMMENT** Contact: Dr. Sergio Verjovski-Almeida  
 Departamento de Bioquimica  
 Instituto de Quimica - Universidade de Sao Paulo  
 Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 Sao Paulo - SP, Brasil  
 Tel: +55-11-3091-2173  
 Fax: +55-11-3091-2186  
 Email: verjov@iq.usp.br  
 This sequence was derived from the FAPESP Schistosoma mansoni EST Genome Project. All sequences in the project were assembled and annotated. This entry and all the assembled sequences can be seen in the following URL <http://bioinfo.iq.usp.br/schisto/>  
 Plate: ML1-0046T-M209 row: 11 column: C.  
 Location/Qualifiers  
 1..441  
 /organism="Schistosoma mansoni"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:6183"  
 /clone="ML1-0046T-M209-C11.G"  
 /sex="mixed pool"  
 /dev\_stage="miracidium"  
 /clone\_1lb="ML1-0046"  
 /note="Vector: pGEM T-easy"



Email: couseens@msu.edu  
Seq primer: M13.

FEATURES  
source  
Location/Qualifiers  
1..221

/organism="Sus scrofa"  
/mol\_type="mRNA"  
/db\_xref="taxon:9823"  
/clone="PBL"  
/sex="male and female"  
/tissue\_type="Brain and central nervous system"  
/dev\_stage="fetal, 10-day, 21-day, 5-week, mature boar,  
post-pubertal gilt, lactating sow"  
/clone\_lib="Porcine Brain Library"  
/note="Porcine (Pig) brain library includes pre-frontal  
cortex, frontal cortex, hippocampus, hypothalamus,  
parietal cortex, amygdala, cerebellum, spinal cord, eye,  
accumbens, and fetal pig CNS"

## ORIGIN

Query Match 80.0%; Score 16.8; DB 14; Length 221;  
Best Local Similarity 90.0%; Pred. No. 1e+03; Indels 0; Gaps 0;  
Matches 18; Conservative 0; Mismatches 2;

QY 2 CGTCGACGCTTCGAGATGAT 21  
|||||  
33 CGTCGACGCTTCGAGATGAT 52

## RESULT 7

CA707068 461 bp mRNA linear EST 26-NOV-2002  
LOCUS CA707068/c  
DEFINITION wdk2c.pk0004.g12 wdk2c Triticum aestivum cDNA clone  
CA707068  
CA707068.1 GI:25428861

EST.  
Triticum aestivum (bread wheat)  
SOURCE  
ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
Poaceae; Triticaceae; Triticum.  
1 (bases 1 to 461)

REFERENCE  
AUTHORS  
Tinney, S.V., Powell, W., Wolters, P., Dolan, M., Hainey, C., Yuan, Z.,  
Miao, G., Garner, N. and Hanafey, M.K.

TITLE  
JOURNAL  
COMMENT  
Unpublished (2002)

CONTACT: Scott V. Tinney  
Crop Genetics  
E. I. Dupont de Nemours and Company  
1 Innovation Way, P.O. Box 6104, Newark, DE 19714-6104, USA  
Tel: 302-631-2602  
Fax: 302-631-2607  
Email: Scott.V.Tinney@usa.dupont.com  
Seq primer: M13.

FEATURES  
source  
Location/Qualifiers  
1..461

/organism="Triticum aestivum"  
/mol\_type="mRNA"  
/db\_xref="taxon:4565"  
/clone="wdk2c.pk0004.g12"  
/tissue\_type="kernel"  
/clone\_lib="wdk2c"  
/note="Vector: pBluescript SK+; Site 1: EcoRI, Site 2:  
XhoI; wheat (Triticum aestivum L.) developing kernel, 7  
days after anthesis."

## ORIGIN

Query Match 80.0%; Score 16.8; DB 14; Length 461;  
Best Local Similarity 90.0%; Pred. No. 1.3e+03; Indels 0; Gaps 0;  
Matches 18; Conservative 0; Mismatches 2;

QY 2 CGTCGACGCTTCGAGATGAT 21  
|||||

DB 293 CGCGACGCTTCGAGATGAT 274

RESULT 8  
CA827269 587 bp mRNA linear EST 11-DEC-2002  
LOCUS CA827269  
DEFINITION 1114012C10.y1 1114 - Unigene IV from Maize Genome Project Zea mays  
CDNA, mRNA sequence.  
CA827269  
CA827269.1 GI:26455686  
EST.  
KEYWORDS  
SOURCE  
ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACAD  
clade; Panicoidae; Andropogoneae; Zea.  
1 (bases 1 to 587)

## REFERENCE

AUTHORS  
TITLE  
JOURNAL  
COMMENT  
Unpublished (1999)  
Contact: Walbot V  
Department of Biological Sciences  
Stanford University  
855 California Ave, Palo Alto, CA 94304, USA  
Tel: 650 723 2227  
Fax: 650 725 8221  
Email: walbot@stanford.edu  
Plate: 1114012 row: C column: 10.

FEATURES  
source  
Location/Qualifiers  
1..587

/organism="Zea mays"  
/mol\_type="mRNA"  
/db\_xref="dbEST:3524.1\_56\_1\_G06.y\_1"  
/db\_xref="taxon:4577"  
/clone\_lib="1114 - Unigene IV from Maize Genome Project"  
/note="This library represents the unique genes found in  
the fourth round of EST sequencing at Stanford University  
for the maize genome project. Sequences are present from  
libraries 1091 and 3524. Contigs were assembled using  
ZmBAssembler and 2 representatives from each contig were  
selected for the Unigene set. All singlets were also  
selected."

## ORIGIN

Query Match 80.0%; Score 16.8; DB 14; Length 587;  
Best Local Similarity 90.0%; Pred. No. 1.4e+03; Indels 0; Gaps 0;  
Matches 18; Conservative 0; Mismatches 2;

QY 1 TCCTCGACGCTTCGAGATGA 20  
|||||  
30 TCCTCGACGCTTCGAGATGA 49

RESULT 9  
BH387860 595 bp DNA linear GSS 11-DEC-2001  
LOCUS BH387860  
DEFINITION AG-ND-129F22.TF ND-TAM Anopheles gambiae genomic clone  
AG-ND-129F22, genomic survey sequence.  
BH387860  
BH387860.1 GI:17334001  
GSS.

ORGANISM  
Anopheles gambiae (African malaria mosquito)  
Anopheles gambiae  
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
Neoptera; Endopterygota; Diptera; Nematocera; Culicoidae;  
Anopheles.

## REFERENCE

AUTHORS  
Hong, Y.S., Hogan, J.R., Wang, X., Sarkar, A., Sim, C., Loftus, B.J.,  
Ren, C., Huff, E.R., Carlile, J.L., Black, K., Zhang, H.-B.,  
Gardner, M.J. and Collins, F.H.  
Construction of a BAC library and generation of BAC end  
sequence-tagged connectors for genome sequencing of the African

JOURNAL  
MEDLINE  
PUBMED  
COMMENT

malaria mosquito Anopheles gambiae  
Mol. Genet. Genomics 268 (6), 720-728 (2003)  
22542063  
12655398  
Other GSSs: AG-ND-129F22, TR  
Contact: Brendan J Loftus  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850, USA  
Tel: 301 838 0208  
Fax: 301 838 3543  
Email: b1o@fugate.igf.org

This clone is from an A. gambiae BAC library (ND-TAM) provided by  
F.H. Collins and sequenced by The Institute for Genomic Research  
(TIGR). The BAC library was generated from A. gambiae PEST strain  
DNA. All DNA was extracted from newly hatched first instar larvae  
to minimize the inclusion of DNA from microorganisms that inhabit  
the gut. The DNA is derived from mixed sexes of larvae. The BAC  
library was constructed at Texas A&M University BAC Center  
University, College Station, Texas 77843-2123, USA using a HindIII  
partial digest.  
Seq primer: M13 For  
Class: BAC ends.

FEATURES  
source  
1. 595  
Location/Qualifiers  
/organism="Anopheles gambiae"  
/mol\_type="genomic DNA"  
/strain="PEST"  
/db\_xref="taxon:7165"  
/clone="AG-ND-129F22"  
/clone\_id="ND-TAM"  
/note="Vector: pECBAC1; Site\_1: HindIII"

ORIGIN

Query Match 80.0%; Score 16.8; DB 28; Length 595;  
Best Local Similarity 90.0%; Pred. No. 1.4e+03;  
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTGAACGTTGAGATGA 20  
|||||  
177 TCGTGAACGTTGAGATGA 158

Db

RESULT 10  
BE052250  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

BE052250 1410 bp mRNA linear EST 08-MAR-2001  
GA\_Ba0035L13f Gossypium arboreum 7-10 dpa fiber library Gossypium  
arboreum cDNA clone GA\_Ba0035L13f, mRNA sequence.  
BE052250  
BE052250.1 GI:8379306  
EST.  
Gossypium arboreum  
Gossypium arboreum  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophytes; Magnoliophyta, eudicotyledons, core eudicots,  
rosids; eurosids II; Malvales; Malvaceae; Malvoideae; Gossypium.  
1 (bases 1 to 1410)  
Wing,R.A., Friesch,D., Yu,Y., Main,D., Rambo,T., Simmons,J.,  
Henry,D., Wood,T.C., Leslie,A. and Wilkins,T.A.  
An integrated analysis of the genetics, development, and evolution  
of the cotton fiber  
Unpublished (2000)  
Contact: Wing RA  
Clemson University Genomics Institute  
Clemson University  
100 Jordan Hall, Clemson, SC 29634, USA  
Tel: 864 656 7288  
Fax: 864 656 4293  
Email: twing@clemson.edu  
Seq primer: TAATACGACTCACTATAGG  
High quality sequence start: 13  
High quality sequence stop: 501.  
Location/Qualifiers

source  
1. 1410  
/organism="Gossypium arboreum"  
/mol\_type="mRNA"  
/strain="AKA"  
/cultivar="8400"  
/db\_xref="taxon:29729"  
/clone="GA\_Ba0035L13f"  
/tissue\_type="Fibers isolated from bolls harvested 7-10  
dpa"  
/lab\_host="E. coli"  
/clone\_id="Gossypium arboreum 7-10 dpa fiber library"  
/note="Vector: pBK-CMV; Site\_1: EcoRI; Site\_2: XhoI"

ORIGIN

Query Match 80.0%; Score 16.8; DB 10; Length 1410;  
Best Local Similarity 94.4%; Pred. No. 1.9e+03;  
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTGAACGTTGAGATGA 20  
|||||  
1356 TCGTGAACGAGCAGATGA 1375

Db

RESULT 11  
AW710594/C  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

AW710594 303 bp mRNA linear EST 25-APR-2000  
e4h01ne.r1 Neurospora crassa evening cDNA library Neurospora crassa  
cDNA clone e4h01ne 3', mRNA sequence.  
AW710594  
AW710594.1 GI:7599686  
EST.  
Neurospora crassa  
Neurospora crassa  
Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;  
Sordariomycetidae; Sordariales; Sordariaceae; Neurospora.  
1 (bases 1 to 303)  
Zhu,H., Lai,H., Kupfer,D., Dunlap,J.C. and Roe,B.A.  
Two Neurospora crassa EST Databases  
Unpublished (1998)  
Contact: Bruce A. Roe, University of Oklahoma, broe@ou.edu  
Department of Chemistry and Biochemistry  
Advanced Center for Genome Technology, University of Oklahoma  
620 Parrington Oval, Norman, OK 73019, USA  
Tel: 405 325 4912  
Fax: 405 325 7762  
Email: broe@ou.edu  
Seq primer: Universal Reverse Primer  
High quality sequence stop: 255.  
Location/Qualifiers  
1. 303  
/organism="Neurospora crassa"  
/mol\_type="mRNA"  
/strain="Strain 30-7 (bd, A)"  
/db\_xref="taxon:5141"  
/clone="e4h01ne"  
/tissue\_type="tissue harvested following 22hr growth in  
dark"  
/clone\_id="Neurospora crassa evening cDNA library"  
/note="Vector: pBluescript SK-, Site\_1: XbaI; Site\_2:  
EcoRI; See: Bell-Pedersen,D., et al. PNAS 93:13096,1996.  
5' end of cDNA cloned into XbaI site of pBluescript; 3'  
end of cDNA cloned into EcoRI site of pBluescript"

ORIGIN

Query Match 78.1%; Score 16.4; DB 10; Length 303;  
Best Local Similarity 94.4%; Pred. No. 1.8e+03;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GTCCAGCGTTGAGATGA 20  
|||||  
232 GTCCAGCGTTGAGATGA 215

Db

```

RESULT 12
LOCUS      BM170465          424 bp      mRNA      linear      EST 04-NOV-2002
DEFINITION BM170465 Nori Satoh unpublished cDNA library, neural complex Clona
            intestinalis cDNA clone rcinc008n07 3', mRNA sequence.
ACCESSION  BM170465
KEYWORDS   EST.
SOURCE     BM170465.1 GI:24560352
ORGANISM   Clona intestinalis
            Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
            Phlebobranchia; Clonidae; Clona.
REFERENCE  1 (bases 1 to 424)
AUTHORS   Satou,Y., Shin-I,T., Kohara,Y. and Satoh,N.
TITLE     Expressed genes in Clona intestinalis (2002c)
JOURNAL   Unpublished (2002)
COMMENT   Contact: Nori Satoh
            Department of Zoology
            Kyoto University
            Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
            Tel: 81-75-753-4081
            Fax: 81-75-705-1113
            Email: satoch@sci.kyoto-u.ac.jp.

FEATURES
            source
            1..424
               /organism="Clona intestinalis"
               /mol_type="mRNA"
               /db_xref="taxon:7719"
               /clone="rcinc008n07"
               /tissue_type="neural complex"
               /clone_lib="Nori Satoh unpublished cDNA library, neural
               complex"

ORIGIN
Query Match      78.1%; Score 16.4; DB 13; Length 424;
Best Local Similarity 94.4%; Pred. No. 2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 TCGTCGACGCTTCGAGAT 18
        |||||
        340 TCGTCGACATTCGAGAT 357

RESULT 13
LOCUS      B1515898          439 bp      mRNA      linear      EST 08-APR-2002
DEFINITION B1515898 B160020B20D02.5 Bee Brain Normalized library, B16 Apis mellifera
            cDNA clone B160020B20D02 5', mRNA sequence.
ACCESSION  B1515898
KEYWORDS   EST.
SOURCE     B1515898.1 GI:15366272
ORGANISM   Apis mellifera (honeybee)
            Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
            Neoptera; Endopterygota; Hymenoptera; Apocrita; Aculeata; Apoidea;
            Apidae; Apis.
REFERENCE  1 (bases 1 to 439)
AUTHORS   Whitfield,C.W., Band,M.R., Bonaldo,M.F., Kumar,C.G., Liu,L.,
            Pardinas,J., Robertson,H.M., Soares,B. and Robinson,G.B.
TITLE     Annotated expressed sequence tags and cDNA microarrays for studies
            of brain and behavior in the honey bee
JOURNAL   Genome Res. 12 (4), 555-566 (2002)
COMMENT   Contact: Gene E. Robinson
            Department of Entomology
            University of Illinois
            505 S. Goodwin Ave., Urbana, IL 61801, USA
            Tel: 217 265 0309
            Fax: 217 244 3499
            Email: generobi@life.uiuc.edu
            This research was funded by the University of Illinois Critical

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Research Initiatives Fund and a Burroughs-Wellcome Trust Innovation
Award in Functional Genomics to G.E. Robinson and an NSF
Postdoctoral Fellowship in Bioinformatics to C.W. Whitfield.
PCR Primers
FORWARD: TAAATGCACTCACTATACGG
BACKWARD: ATTATCCCTCACTAAG
Plate: B160020B20 row: D column: 02
Seq primer: AGCGATACATTCACACAGGA
High quality sequence stop: 439.

FEATURES
            source
            1..439
               /organism="Apis mellifera"
               /mol_type="mRNA"
               /strain="mixed strains of European bees, predominantly
               A.m. ligustica"
               /db_xref="taxon:7460"
               /clone="B160020B20D02"
               /sex="female"
               /tissue_type="brain"
               /dev_stage="adult worker honey bee"
               /lab_host="DH10B"
               /clone_lib="Bee Brain Normalized library, B16"
               /note="Organ: brain; Vector: pT73-Pac; Site: 1; EcoRI;
               Site: 2; NotI; The B16 library was contributed by the
               Soares laboratory and it was constructed and normalized
               as described by Bonaldo, M.F., Lennon, G. and Soares,
               M.B. (1996), Genome Research 6(9): 791-806. RNA was
               prepared from dissected brains of adult worker bees of
               various ages and various behavioral groups."

ORIGIN
Query Match      78.1%; Score 16.4; DB 12; Length 439;
Best Local Similarity 94.4%; Pred. No. 2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      4 TCGAAGCTTCGAGATGAT 21
        |||||
        269 TCGAAGCTTCGAGTTGAT 286

RESULT 14
LOCUS      BM300285/c          445 bp      mRNA      linear      EST 11-NOV-2002
DEFINITION BM300285 Nori Satoh unpublished cDNA library, neural complex Clona
            intestinalis cDNA clone cinc008n07 5', mRNA sequence.
ACCESSION  BM300285
KEYWORDS   EST.
SOURCE     BM300285.1 GI:24880896
ORGANISM   Clona intestinalis
            Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
            Phlebobranchia; Clonidae; Clona.
REFERENCE  1 (bases 1 to 445)
AUTHORS   Satou,Y., Shin-I,T., Kohara,Y. and Satoh,N.
TITLE     Expressed genes in Clona intestinalis (2002c)
JOURNAL   Unpublished (2002)
COMMENT   Contact: Nori Satoh
            Department of Zoology
            Kyoto University
            Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
            Tel: 81-75-753-4081
            Fax: 81-75-705-1113
            Email: satoch@sci.kyoto-u.ac.jp.

FEATURES
            source
            1..445
               /organism="Clona intestinalis"
               /mol_type="mRNA"
               /db_xref="taxon:7719"
               /clone="cinc008n07"
               /tissue_type="neural complex"
               /clone_lib="Nori Satoh unpublished cDNA library, neural
               complex"

ORIGIN

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Query Match 78.1%; Score 16.4; DB 13; Length 445;  
 Best Local Similarity 94.4%; Pred. No. 2e+03;  
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGAACGTTCCGAGAT 18  
 |||||  
 Db 80 TCGTCGAACATTCGAGAT 63

RESULT 15  
 BP018564/c 457 bp mRNA linear EST 15-MAR-2002  
 LOCUS BP018564 Nori Satoh unpublished cDNA library, young adult Ciona  
 DEFINITION Bp018564 Nori Satoh unpublished cDNA library, young adult Ciona  
 accession Bp018564  
 VERSION Bp018564  
 KEYWORDS Bp018564.1 GI:19510514  
 EST.  
 SOURCE Ciona intestinalis  
 ORGANISM Ciona intestinalis  
 Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;  
 Phlebobranchia; Clonidae; Ciona.  
 1 (bases 1 to 457)  
 Satoh, N., Satou, Y., Kohara, Y. and Shin-I, T.  
 Expressed genes in Ciona intestinalis  
 Unpublished (2000)  
 CONTACT: Nori Satoh  
 Department of Zoology  
 Kyoto University  
 Sakyo-ku, Kyoto, Kyoto 606-8502, Japan  
 Tel: 81-75-753-4081  
 Fax: 81-75-705-1113  
 Email: satoh@ascidian.zool.kyoto-u.ac.jp.  
 Location/Qualifiers  
 1..457  
 /organism="Ciona intestinalis"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:7719"  
 /clone="clad54123"  
 /tissue\_type="whole animal"  
 /dev\_stage="young adult"  
 /clone\_lib="Nori Satoh unpublished cDNA library, young adult"

ORIGIN  
 Query Match 78.1%; Score 16.4; DB 12; Length 457;  
 Best Local Similarity 94.4%; Pred. No. 2e+03;  
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGAACGTTCCGAGAT 18  
 |||||  
 Db 348 TCGTCGAACATTCGAGAT 331

RESULT 16  
 B29433 504 bp DNA linear GSS 13-OCT-1997  
 LOCUS B29433 F19L1TFB IGF Arabidopsis thaliana genomic clone F19L12, genomic  
 DEFINITION survey sequence.  
 accession B29433  
 VERSION B29433  
 KEYWORDS B29433.1 GI:2515399  
 GSS.  
 SOURCE Arabidopsis thaliana (chale cress)  
 ORGANISM Arabidopsis thaliana  
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
 rosoids; eurosids II; Brassicales; Brassicaceae; Arabidopsi  
 1 (bases 1 to 504)  
 Rounsley, S.D., Kelley, J.M., Field, C.E., Craven, M.B., Adams, M.D. and  
 Venter, J.C.  
 Use of a BAC End Sequence Database to Identify Minimal Overlaps for  
 Arabidopsis Genomic Sequencing  
 Unpublished (1997)

COMMENT  
 Other GSSs: F19L12R  
 Contact: Steve Rounsley  
 Department of Eukaryotic Genomics  
 The Institute for Genomic Research  
 9712 Medical Center Dr., Rockville, MD 20850, USA  
 Tel: 301 838 0200  
 Fax: 301 838 0208  
 Email: rounsley@igr.org  
 Seq primer: M13-21  
 Class: BAC ends  
 High quality sequence stop: 504.  
 Location/Qualifiers  
 1..504  
 /organism="Arabidopsis thaliana"  
 /mol\_type="genomic DNA"  
 /strain="Columbia"  
 /db\_xref="taxon:3702"  
 /clone="F19L12"  
 /sex="hermaphrodite"  
 /clone\_lib="IGF"  
 /note="Vector: BelosACII, Site\_1: EcoRI, Site\_2: EcoRI;  
 Produced by Thomas Altman"

ORIGIN  
 Query Match 78.1%; Score 16.4; DB 28; Length 504;  
 Best Local Similarity 94.4%; Pred. No. 2.1e+03;  
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 TCGAACGTTCCGAGAT 21  
 |||||  
 Db 359 TCGATCGTTCCGAGAT 342

RESULT 17  
 B1514876 511 bp mRNA linear EST 08-APR-2002  
 LOCUS B1514876 Bb160016B10G01.5 Bee Brain Normalized library, Bb16 Apis mellifera  
 DEFINITION cDNA clone Bb160016B10G01.5', mRNA sequence.  
 accession B1514876  
 VERSION B1514876  
 KEYWORDS B1514876.1 GI:15365250  
 EST.  
 SOURCE Apis mellifera (honeybee)  
 ORGANISM Apis mellifera  
 Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
 Neoptera; Endopterygota; Hymenoptera; Apocrita; Aculeata; Apoidea;  
 Apidae; Apis.  
 1 (bases 1 to 511)  
 Whitfield, C.W., Band, M.R., Bonaldi, M.F., Kumar, C.G., Liu, L.,  
 Pardinas, J., Robertson, H.M., Soares, B. and Robinson, G.E.  
 Annotated expressed sequence tags and cDNA microarrays for studies  
 of brain and behavior in the honey bee  
 Genome Res. 12 (4), 555-566 (2002)  
 MEDLINE 21929762  
 PUBMED 11932240  
 COMMENT Contact: Gene E. Robinson  
 Department of Entomology  
 University of Illinois  
 505 S. Goodwin Ave., Urbana, IL 61801, USA  
 Tel: 217 265 0309  
 Fax: 217 244 3499  
 Email: generobi@life.uiuc.edu  
 This research was funded by the University of Illinois Critical  
 Research Initiatives Fund and a Burroughs-Wellcome Trust Innovation  
 Award in Functional Genomics and a G.E. Robinson and an NSF  
 Postdoctoral Fellowship in Bioinformatics to C.W. Whitfield.  
 PCR primers  
 FORWARD: TTAATGACCTCAGATAGG  
 BACKWARD: ATTATCCCTCAGTAAG  
 Plate: Bb160016B10 row: G column: 01  
 Seq primer: AGCGATACACATTCACACAGCA  
 High quality sequence stop: 511.  
 Location/Qualifiers  
 1..511

FEATURES  
 SOURCE

/organism="Apis mellifera"  
/mol\_type="mRNA"  
/strain="mixed strains of European bees, predominantly  
A.m. ligustica"  
/db\_xref="taxon:7460"  
/clone="BB160016B10G01"  
/sex="female"  
/tissue\_type="brain"  
/dev\_stage="adult worker honey bee"  
/lab\_host="DH10B"  
/clone\_lib="Bee Brain Normalized Library, BB16"  
/note="Organ: brain; Vector: pT73-Pac; Site 1: Ecot1;  
Site 2: Not1; The BB16 library was contributed by the  
Soares laboratory and it was constructed and normalized  
as described by Bonaldo, M.F., Lennon, G. and Soares,  
M.B. (1996), Genome Research 6(9): 791-806. RNA was  
prepared from dissected brains of adult worker bees of  
various ages and various behavioral groups."

## .ORIGIN

Query Match 78.1%; Score 16.4; DB 12; Length 511;  
Best Local Similarity 94.4%; Pred. No. 2.1e+03;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 TCGAACGTTTCGAGATGAT 21  
|||||  
Db 401 TCGAACGTTTCGAGATGAT 418

RESULT 18 511 bp mRNA linear EST 08-APR-2002  
LOCUS BB1514893  
DEFINITION BB160016B10H09.5 Bee Brain Normalized Library, BB16 Apis mellifera  
CDNA clone BB160016B10H09.5, mRNA sequence.  
ACCESSION BB1514893  
VERSION BB1514893.1 GI:15365267  
KEYWORDS EST.  
SOURCE Apis mellifera (honeybee)  
ORGANISM Apis mellifera

REFERENCE 1 (bases 1 to 511)  
AUTHORS Whitfield, C.W., Band, M.R., Bonaldo, M.F., Kumar, C.G., Liu, L.,  
Parinaua, J., Robertson, H.M., Soares, B., and Robinson, G.E.  
TITLE Annotated expressed sequence tags and cDNA microarrays for studies  
of brain and behavior in the honey bee  
JOURNAL Genome Res. 12 (4), 555-566 (2002)  
MEDLINE 21929762  
PUBMED 11932240

COMMENT Contact: Gene E. Robinson  
Department of Entomology  
University of Illinois  
505 S. Goodwin Ave., Urbana, IL 61801, USA  
Tel: 217 265 0309  
Fax: 217 244 3499

Email: generobi@life.uiuc.edu  
This research was funded by the University of Illinois Critical  
Research Initiatives Fund and a Burroughs-Wellcome Trust Innovation  
Award in Functional Genomics to G.E. Robinson and an NSF  
Postdoctoral Fellowship in Bioinformatics to C.W. Whitfield.  
PCR Primers  
FORWARD: TAAATGACTCACTATAGG  
BACKWARD: ATTAACCTCACTAAG  
Plate: BB160016B10 row: H column: 09  
Seq primer: AGCGATACATTCACACAGCA  
High quality sequence stop: 511.  
location/Qualifiers

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1..511  
/organism="Apis mellifera"  
/mol\_type="mRNA"  
/strain="mixed strains of European bees, predominantly  
A.m. ligustica"

/db\_xref="taxon:7460"  
/clone="BB160016B10H09"  
/sex="female"  
/tissue\_type="brain"  
/dev\_stage="adult worker honey bee"  
/lab\_host="DH10B"  
/clone\_lib="Bee Brain Normalized Library, BB16"  
/note="Organ: brain; Vector: pT73-Pac; Site 1: Ecot1;  
Site 2: Not1; The BB16 library was contributed by the  
Soares laboratory and it was constructed and normalized  
as described by Bonaldo, M.F., Lennon, G. and Soares,  
M.B. (1996), Genome Research 6(9): 791-806. RNA was  
prepared from dissected brains of adult worker bees of  
various ages and various behavioral groups."

## .ORIGIN

Query Match 78.1%; Score 16.4; DB 12; Length 511;  
Best Local Similarity 94.4%; Pred. No. 2.1e+03;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 TCGAACGTTTCGAGATGAT 21  
|||||  
Db 401 TCGAACGTTTCGAGATGAT 418

RESULT 19 635 bp mRNA linear EST 24-OCT-2002  
LOCUS BB118728  
DEFINITION BB118728 Nori Satoh unpublished cDNA library, tailbud embryo Clona  
intestinalis cDNA clone rcitb080n02 3', mRNA sequence.  
ACCESSION BB118728  
VERSION BB118728.1 GI:24365393  
KEYWORDS EST.  
SOURCE Clona intestinalis  
ORGANISM Clona intestinalis

REFERENCE 1 (bases 1 to 635)  
AUTHORS Satou, Y., Shin-i, T., Kohara, Y. and Satoh, N.  
TITLE Expressed genes in Clona intestinalis (2002)  
JOURNAL Unpublished (2002)  
COMMENT Contact: Nori Satoh  
Department of Zoology  
Kyoto University  
Sakyo-ku, Kyoto 606-8502, Japan  
Tel: 81-75-753-4081  
Fax: 81-75-705-1113  
Email: satoh@scidian.zool.kyoto-u.ac.jp.  
location/Qualifiers

## FEATURES

1..635  
/organism="Clona intestinalis"  
/mol\_type="mRNA"  
/db\_xref="taxon:7719"  
/clone="rcitb080n02"  
/tissue\_type="whole animal"  
/dev\_stage="tailbud embryo"  
/clone\_lib="Nori Satoh unpublished cDNA library, tailbud  
embryo"

## .ORIGIN

Query Match 78.1%; Score 16.4; DB 13; Length 635;  
Best Local Similarity 94.4%; Pred. No. 3.3e+03;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGACGTTTCGAGAT 18  
|||||  
Db 331 TCGTCGACGTTTCGAGAT 348

RESULT 20 650 bp mRNA linear EST 09-NOV-2001  
LOCUS AV902415  
DEFINITION AV902415 Nori Satoh unpublished cDNA library, young adult Clona



intestinalis cDNA clone rc1ad54123 3', mRNA sequence.

AV902415 GI:16891513

AV902415.1

EST.

Clona intestinalis

ORGANISM Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona; Phlebobranchia; Cloniidae; Clona.

REFERENCE 1 (bases 1 to 650)

AUTHORS Satoh,N., Satou,Y., Kohara,Y. and Shin-I,T.

TITLE Expressed genes in Clona intestinalis

JOURNAL Unpublished (2000)

COMMENT Contact: Nori Satoh

Department of Zoology

Kyoto University

Sakyo-ku, Kyoto, Kyoto 606-8502, Japan

Tel: 81-75-753-4081

Fax: 81-75-705-1113

Email: satoh@ascidian.zool.kyoto-u.ac.jp.

Location/Qualifiers

1..650

/organism="Clona intestinalis"

/mol\_type="mRNA"

/db\_xref="taxon:7719"

/clone="rc1ad54123"

/tissue\_type="whole animal"

/dev\_stage="young adult"

/clone\_lib="Nori Satoh unpublished cDNA library, young adult"

ORIGIN

Query Match 78.1%; Score 16.4; DB 9; Length 650;

Best Local Similarity 94.4%; Pred. No. 2.3e+03;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGAACGTTCCGAGT 18

Db 349 TCGTCGAACATTCGAGAT 366

RESULT 21

BM174317 662 bp mRNA linear EST 04-NOV-2002

LOCUS BM174317

DEFINITION BM174317 Nori Satoh unpublished cDNA library, neural complex Clona intestinalis cDNA clone rcinc030b19 3', mRNA sequence.

BM174317

ACCESSION BM174317.1 GI:24564241

VERSION EST.

KEYWORDS Clona intestinalis

SOURCE Clona intestinalis

ORGANISM Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona; Phlebobranchia; Cloniidae; Clona.

REFERENCE 1 (bases 1 to 662)

AUTHORS Satou,Y., Shin-I,T., Kohara,Y. and Satoh,N.

TITLE Expressed genes in Clona intestinalis (2002c)

JOURNAL Unpublished (2002)

COMMENT Contact: Nori Satoh

Department of Zoology

Kyoto University

Sakyo-ku, Kyoto, Kyoto 606-8502, Japan

Tel: 81-75-753-4081

Fax: 81-75-705-1113

Email: satoh@ascidian.zool.kyoto-u.ac.jp.

Location/Qualifiers

1..662

/organism="Clona intestinalis"

/mol\_type="mRNA"

/db\_xref="taxon:7719"

/clone="rcinc030b19"

/tissue\_type="neural complex"

/clone\_lib="Nori Satoh unpublished cDNA library, neural complex"

ORIGIN

Query Match 78.1%; Score 16.4; DB 13; Length 662;

Best Local Similarity 94.4%; Pred. No. 2.3e+03;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGAACGTTCCGAGT 18

Db 328 TCGTCGAACATTCGAGAT 345

RESULT 22

BM171245 663 bp mRNA linear EST 04-NOV-2002

LOCUS BM171245

DEFINITION BM171245 Nori Satoh unpublished cDNA library, neural complex Clona intestinalis cDNA clone rcinc018d19 3', mRNA sequence.

BM171245

ACCESSION BM171245.1 GI:24561132

VERSION EST.

KEYWORDS Clona intestinalis

SOURCE Clona intestinalis

ORGANISM Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona; Phlebobranchia; Cloniidae; Clona.

REFERENCE 1 (bases 1 to 663)

AUTHORS Satou,Y., Shin-I,T., Kohara,Y. and Satoh,N.

TITLE Expressed genes in Clona intestinalis (2002c)

JOURNAL Unpublished (2002)

COMMENT Contact: Nori Satoh

Department of Zoology

Kyoto University

Sakyo-ku, Kyoto, Kyoto 606-8502, Japan

Tel: 81-75-753-4081

Fax: 81-75-705-1113

Email: satoh@ascidian.zool.kyoto-u.ac.jp.

Location/Qualifiers

1..663

/organism="Clona intestinalis"

/mol\_type="mRNA"

/db\_xref="taxon:7719"

/clone="rcinc018d19"

/tissue\_type="neural complex"

/clone\_lib="Nori Satoh unpublished cDNA library, neural complex"

ORIGIN

Query Match 78.1%; Score 16.4; DB 13; Length 663;

Best Local Similarity 94.4%; Pred. No. 2.3e+03;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGAACGTTCCGAGT 18

Db 329 TCGTCGAACATTCGAGAT 346

RESULT 23

BM297679/c 707 bp mRNA linear EST 11-NOV-2002

LOCUS BM297679

DEFINITION BM297679 Nori Satoh unpublished cDNA library, neural complex Clona intestinalis cDNA clone rcinc018d19 5', mRNA sequence.

BM297679

ACCESSION BM297679.1 GI:24878290

VERSION EST.

KEYWORDS Clona intestinalis

SOURCE Clona intestinalis

ORGANISM Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona; Phlebobranchia; Cloniidae; Clona.

REFERENCE 1 (bases 1 to 707)

AUTHORS Satou,Y., Shin-I,T., Kohara,Y. and Satoh,N.

TITLE Expressed genes in Clona intestinalis (2002c)

JOURNAL Unpublished (2002)

COMMENT Contact: Nori Satoh

Department of Zoology

Kyoto University

Sakyo-ku, Kyoto, Kyoto 606-8502, Japan

Tel: 81-75-753-4081  
Fax: 81-75-705-1113  
Email: satcho@ascidian.zool.kyoto-u.ac.jp.

## FEATURES

source Location/Qualifiers

1.707  
/organism="Ciona intestinalis"  
/mol\_type="mRNA"  
/db\_xref="taxon:7719"  
/clone="cinc018d19"  
/tissue\_type="neural complex"  
/clone\_1ib="Nori Satcho unpublished cDNA library, neural complex"

## ORIGIN

Query Match 78.1%; Score 16.4; DB 13; Length 707;  
Best Local Similarity 94.4%; Pred. No. 2.3e+03;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCCTCGACGCTCGAGAT 18  
|||||  
Db 342 TCCTCGACATTCGAGAT 325

RESULT 24 709 bp mRNA linear EST 11-NOV-2002  
BM303599/c  
LOCUS BM303599 Nori Satcho unpublished cDNA library, neural complex Ciona  
DEFINITION  
BM303599  
Accession  
BM303599.1 GI:24884210  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM

Ciona intestinalis  
Ciona intestinalis  
Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;  
Phlebobranchia; Clonidae; Ciona.

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT  
1 (bases 1 to 709)  
Satcho, Y., Shin-I, T., Kohara, Y. and Satcho, N.  
Expressed genes in Ciona intestinalis (2002c)  
Unpublished (2002)  
Contact: Nori Satcho  
Department of Zoology  
Kyoto University  
Sakyo-ku, Kyoto 606-8502, Japan  
Tel: 81-75-753-4081  
Fax: 81-75-705-1113  
Email: satcho@ascidian.zool.kyoto-u.ac.jp.

## FEATURES

source Location/Qualifiers

1.709  
/organism="Ciona intestinalis"  
/mol\_type="mRNA"  
/db\_xref="taxon:7719"  
/clone="cinc030d19"  
/tissue\_type="neural complex"  
/clone\_1ib="Nori Satcho unpublished cDNA library, neural complex"

## ORIGIN

Query Match 78.1%; Score 16.4; DB 13; Length 709;  
Best Local Similarity 94.4%; Pred. No. 2.4e+03;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCCTCGACGCTCGAGAT 18  
|||||  
Db 344 TCCTCGACATTCGAGAT 327

RESULT 25 726 bp mRNA linear EST 04-NOV-2002  
BM174467  
LOCUS BM174467 Nori Satcho unpublished cDNA library, neural complex Ciona  
DEFINITION  
BM174467  
Accession  
BM174467.1 GI:24564391  
VERSION

KEYWORDS  
SOURCE  
ORGANISM  
Ciona intestinalis  
Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;  
Phlebobranchia; Clonidae; Ciona.

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT  
1 (bases 1 to 726)  
Satcho, Y., Shin-I, T., Kohara, Y. and Satcho, N.  
Expressed genes in Ciona intestinalis (2002c)  
Unpublished (2002)  
Contact: Nori Satcho  
Department of Zoology  
Kyoto University  
Sakyo-ku, Kyoto 606-8502, Japan  
Tel: 81-75-753-4081  
Fax: 81-75-705-1113  
Email: satcho@ascidian.zool.kyoto-u.ac.jp.

## FEATURES

source

1.726  
/organism="Ciona intestinalis"  
/mol\_type="mRNA"  
/db\_xref="taxon:7719"  
/clone="rcinc030j13"  
/tissue\_type="neural complex"  
/clone\_1ib="Nori Satcho unpublished cDNA library, neural complex"

## ORIGIN

Query Match 78.1%; Score 16.4; DB 13; Length 726;  
Best Local Similarity 94.4%; Pred. No. 2.4e+03;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCCTCGACGCTCGAGAT 18  
|||||  
Db 329 TCCTCGACATTCGAGAT 346

RESULT 26 914 bp DNA linear GSS 01-SEP-2000  
CNS04A00  
LOCUS CNS04A00  
DEFINITION  
Tetradon nigroviridis genome survey sequence PUC-Orl end of clone  
095014 of library G from Tetradon nigroviridis, genomic survey  
sequence.

ACCESSION  
AL282105  
VERSION  
AL282105.1 GI:8020441  
KEYWORDS  
GSS: genome survey sequence.  
SOURCE  
Tetradon nigroviridis  
ORGANISM  
Tetradon nigroviridis

REFERENCE  
AUTHORS  
TITLE  
1 Roest Crolius, H., Jallion, O., Dasilva, C., Bouneau, L., Fisher, C., Bernot, A., Fizames, C., Wincker, P., Brotlier, P., Quetier, F., Saurin, W. and Weissenbach, J.  
Estimate of human gene number provided by genome-wide analysis using Tetradon nigroviridis DNA sequence  
Nat. Genet. 25 (2), 235-238 (2000)

REFERENCE  
AUTHORS  
TITLE  
2 Roest Crolius, H., Jallion, O., Dasilva, C., Ozouf-Costaz, C., Fizames, C., Fischer, C., Bouneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A. and Weissenbach, J.  
Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetradon nigroviridis  
Genome Res. 10 (7), 939-949 (2000)

JOURNAL  
MEDLINE  
PUBMED  
20296633  
10835645

## TITLE

## JOURNAL

## MEDLINE

## PUBMED

## AUTHORS

## TITLE

## JOURNAL

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## PUBMED

## AUTHORS

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## JOURNAL

## PUBMED

## AUTHORS

## COMMENT

BP 191 91006 EVRY cedex - FRANCE (E-mail : segref@genoscope.cns.fr  
- web : www.genoscope.cns.fr)  
This sequence is a single read and was generated as part of a large  
scale clone-end sequencing project of the Tetraodon nigroviridis  
genome. For more information, please take a look at  
<http://www.genoscope.cns.fr/Tetraodon>.

## FEATURES

## SOURCE

1..914  
/organism="Tetraodon nigroviridis"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:99883"  
/clone\_id="095j14"  
/note="Genoscope sequence ID : C08G095DE07SP1-end ;  
PUC-Or1"

## ORIGIN

Query Match 78.1% Score 16.4; DB 29; Length 914;  
Best Local Similarity 94.4%; Pred. No. 2.5e+03;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 GTCCAGCCTTCGAGATGA 20  
|||||  
333 GTCCAGCCTTCAGATGA 350

Db 333 GTCCAGCCTTCAGATGA 350

RESULT 27 166 bp mRNA linear EST 25-SEP-2003  
CA196886  
LOCUS  
DEFINITION SCFPAD1093H09.g AD1 Saccharum officinarum cDNA clone SCBPAD1093H09  
5' mRNA sequence.

ACCESSION CA196886  
VERSION CA196886.1 GI:35226590  
KEYWORDS EST.  
SOURCE Saccharum officinarum  
ORGANISM Saccharum officinarum

REFERENCE Eukaryota, Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD  
clade; Panicoideae; Andropogoneae; Saccharum.  
1 (bases 1 to 166)  
Vettore,A.L., da Silva,F.R., Kemper,E.L. and Arruda,P.  
The libraries that made SUCEST  
Genet. Mol. Biol. 24 (1-4), 1-7 (2001)

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## COMMENT

Contact: Arruda P  
Centro de Biologia Molecular e Engenharia Genetica  
Universidade Estadual de Campinas  
Caixa Postal 6010, 13083-970, Campinas SP, Brazil  
Tel: 55 19 3788 1137  
Fax: 55 19 3788 1089  
Email: patricia@unicamp.br

Clone distribution: clone distribution information can be found  
through the Brazilian Clone Collection Center (BCCC) at  
<http://www.bcccenter.fcav.unesp.br>  
Plate: 093 row: H column: 09  
Seq primer: T7 Promoter Primer.  
Location/Qualifiers

## FEATURES

## SOURCE

1..166  
/organism="Saccharum officinarum"  
/mol\_type="mRNA"  
/db\_xref="taxon:4547"  
/clone\_id="SCFPAD1093H09"  
/lab\_host="DH10B"  
/clone\_id="AD1"  
/note="Organ: seedlings inoculated with Gluconacetobacter  
diazotrophicus; Vector: pSport1; Site 1: SalI; Site 2:  
NotI; An unidirectional cDNA library generated from  
[seedlings inoculated with Gluconacetobacter  
diazotrophicus]. cDNA was prepared from polyA+ mRNA using  
SuperScript Plasmid System Kit (Invitrogen). The  
double-strand cDNAs were fractionated in a sepharose  
CL-2B 40cm-columns and fragments sizing between 0.8 and  
1.5 Kb were directionally cloned into the vector. Details  
of each source of RNA and library construction can be

## ORIGIN

obtained at <http://sucest.lad.ic.unicamp.br/public>

Query Match 77.1% Score 16.2; DB 14; Length 166;  
Best Local Similarity 85.7%; Pred. No. 1.9e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 TCGTCGACGTTTCGAGATGAT 21  
|||||  
106 TCGTCGACGTTTCGAGATGAT 126

Db 106 TCGTCGACGTTTCGAGATGAT 126

RESULT 28  
BG931701/c

LOCUS BG931701 273 bp mRNA linear EST 03-JUL-2002  
DEFINITION h112-118 h112 S. mansoni adult mini-library,  
Fietto/Demarco/Verjovski-Almeida Schistosoma mansoni cDNA, mRNA  
sequence.

ACCESSION BG931701.1 GI:17156638  
VERSION BG931701.1  
KEYWORDS EST.  
SOURCE Schistosoma mansoni  
ORGANISM Schistosoma mansoni

REFERENCE Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;  
Strigeidida; Schistosomatoidae; Schistosomatidae; Schistosoma.  
1 (bases 1 to 273)  
Fietto,J.U.R., Demarco,R. and Verjovski-Almeida,S.  
Use of degenerate primers and touchdown PCR for construction of  
cDNA libraries  
Biotechniques 32 (6), 1404-1408 (2002)

JOURNAL MEDLINE  
PUBMED 12074173  
Contact: Verjovski-Almeida S  
Departamento de Bioquímica, Instituto de Química  
Universidade de São Paulo  
Av. Lineu Prestes, 748, São Paulo, SP 05508-900, Brasil  
Tel: 55-11-3091-2173  
Fax: 55-11-3091-2186  
Email: verjovski@usp.br

COMMENT  
PUBMED 12074173  
Contact: Verjovski-Almeida S  
Departamento de Bioquímica, Instituto de Química  
Universidade de São Paulo  
Av. Lineu Prestes, 748, São Paulo, SP 05508-900, Brasil  
Tel: 55-11-3091-2173  
Fax: 55-11-3091-2186  
Email: verjovski@usp.br

## FEATURES

## SOURCE

1..273  
/organism="Schistosoma mansoni"  
/mol\_type="mRNA"  
/strain="B1"  
/db\_xref="taxon:6183"  
/dev\_stage="Adult"  
/clone\_id="h112 S. mansoni adult mini-library,  
Fietto/Demarco/Verjovski-Almeida"  
/note="Vector: Bluescript SK; minilibrary constructed  
using low-stringency RT-PCR and consensus-degenerate  
primer"

## ORIGIN

Query Match 77.1% Score 16.2; DB 12; Length 273;  
Best Local Similarity 85.7%; Pred. No. 2.2e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 TCGTCGACGTTTCGAGATGAT 21  
|||||  
250 TCGTCGACGTTTCGAGATGAT 230

Db 250 TCGTCGACGTTTCGAGATGAT 230

RESULT 29  
CD190559/c

LOCUS MS1-0064U-A263-G08-U.B MS1-0064 Schistosoma mansoni cDNA clone  
DEFINITION MS1-0064U-A263-G08-B, mRNA sequence.  
ACCESSION CD190559  
VERSION CD190559.1 GI:34720529  
KEYWORDS EST.  
SOURCE Schistosoma mansoni  
ORGANISM Schistosoma mansoni

Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
COMMENT

Strigoidae; Schistosomatidae; Schistosomatidae; Schistosoma.  
1 (bases 1 to 314)  
Verjovski-Almeida, S., Demarco, R., Martins, E.A.L., Guimaraes, P.E.M.,  
Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y. Jr.,  
Kiteajima, U.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F.,  
Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L.,  
Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A.,  
Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A.,  
Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T.,  
Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M.,  
Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.  
Transcriptome analysis of the acclimated human parasite Schistosoma  
mansoni  
Nat. Genet. 35 (2), 148-157 (2003)  
22879926  
Contact: Dr. Sergio Verjovski-Almeida  
Departamento de Bioquímica  
Instituto de Química - Universidade de São Paulo  
Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 São Paulo - SP,  
Brasil  
Tel: +55-11-3091-2173  
Fax: +55-11-3091-2186  
Email: verjo@iq.usp.br  
This sequence was derived from the FAPESP Schistosoma mansoni EST  
Genome Project. All sequences in the project were assembled and  
annotated. This entry and all the assembled sequences can be seen  
in the following URL: <http://bioinfo.iq.usp.br/schisto/>  
Plate: MS1-0064U-A263 row: 8 column: G.

FEATURES  
source  
1..314  
/organism="Schistosoma mansoni"  
/mol\_type="mRNA"  
/db\_xref="taxon:6183"  
/clone="MS1-0064U-A263-G08.B"  
/sex="mixed pool"  
/dev\_stage="schistosomulum"  
/lab\_host="in vitro culture"  
/clone\_lib="MS1-0064"  
/note="Vector: pGEM T-easy"

ORIGIN  
Query Match 77.1%; Score 16.2; DB 14; Length 314;  
Best Local Similarity 85.7%; Pred. No. 2.3e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY  
1 TCGTCGACGTTGAGATGAT 21  
|||||  
65 TCGCTCACTTCGACATGAT 45

Db  
315 bp mRNA linear EST 14-SEP-2003  
CD183371  
MS1-0038U-A246-C02-U.B MS1-0038 Schistosoma mansoni cDNA clone  
CD183371  
MS1-0038U-A246-C02.B, mRNA sequence.

ACCESSION  
CD183371  
GI:34713578

VERSION  
EST

KEYWORDS  
Schistosoma mansoni  
Schistosoma mansoni  
Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;  
Strigoidae; Schistosomatidae; Schistosomatidae; Schistosoma.

SOURCE  
1 (bases 1 to 315)  
Verjovski-Almeida, S., Demarco, R., Martins, E.A.L., Guimaraes, P.E.M.,  
Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y. Jr.,  
Kiteajima, U.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F.,  
Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L.,  
Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A.,  
Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A.,  
Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T.,  
Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M.,  
Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.  
Transcriptome analysis of the acclimated human parasite Schistosoma

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
COMMENT

mansoni  
Nat. Genet. 35 (2), 148-157 (2003)  
22879926  
Contact: Dr. Sergio Verjovski-Almeida  
Departamento de Bioquímica  
Instituto de Química - Universidade de São Paulo  
Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 São Paulo - SP,  
Brasil  
Tel: +55-11-3091-2173  
Fax: +55-11-3091-2186  
Email: verjo@iq.usp.br  
This sequence was derived from the FAPESP Schistosoma mansoni EST  
Genome Project. All sequences in the project were assembled and  
annotated. This entry and all the assembled sequences can be seen  
in the following URL: <http://bioinfo.iq.usp.br/schisto/>  
Plate: MS1-0038U-A246 row: 2 column: C.

FEATURES  
source  
1..315  
/organism="Schistosoma mansoni"  
/mol\_type="mRNA"  
/db\_xref="taxon:6183"  
/clone="MS1-0038U-A246-C02.B"  
/sex="mixed pool"  
/dev\_stage="schistosomulum"  
/lab\_host="in vitro culture"  
/clone\_lib="MS1-0038"  
/note="Vector: pGEM T-easy"

ORIGIN  
Query Match 77.1%; Score 16.2; DB 14; Length 315;  
Best Local Similarity 85.7%; Pred. No. 2.3e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY  
1 TCGTCGACGTTGAGATGAT 21  
|||||  
248 TCGCTCACTTCGACATGAT 268

Db  
328 bp mRNA linear EST 20-JUN-2001  
BI075646  
IP1\_23\_C09.g1 A002 Immature pannicle 1 (IP1) Sorghum bicolor cDNA,  
mRNA sequence.

ACCESSION  
BI075646  
GI:14514303

VERSION  
EST

KEYWORDS  
Sorghum bicolor (sorghum)  
Sorghum bicolor  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD  
clade; Panicoidae; Andropogoneae; Sorghum.

SOURCE  
1 (bases 1 to 328)  
Klein, R.R., Cordonnier-Pratt, M.-M., Gingle, A., Sudman, M. and  
Pratt, L.H.  
An EST database from Sorghum: developing preanthesis pannicles  
Unpublished (2001)  
Contact: Cordonnier-Pratt MM  
Laboratory for Genomics and Bioinformatics  
The University of Georgia, Department of Plant Biology  
Plant Sciences Building, Km. 2502, Athens, GA 30602-7271, USA  
Tel: 706 542 1860  
Fax: 706 583 0210  
Email: mmprratt@uga.edu  
Sequences have been trimmed to exclude polyA, vector and regions  
below phred quality 16. The threshold for high quality sequence is  
20. Three-prime sequences, which are obtained with PolyTm1x or T7  
sequencing primer, are presented as the reverse complement.

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
COMMENT

High quality sequence start: 4  
High quality sequence stop: 328  
POLYA=yes.  
Location/Qualifiers  
1..328

/organism="Sorghum bicolor"  
/mol\_type="mRNA"  
/cultivar="Brix623"  
/db\_xref="taxon:4558"  
/clone\_lib="Immature panicle 1 (IP1)"  
/note="Organ: Developing preanthesis panicles; Vector: pBluescript II SK(-) from Lambda Zap II; Site 1: XhoI; Site 2: EcoRI; The library was made from poly-A RNA in the cloning vector Lambda Zap II. Clones to be sequenced were prepared by mass excision."

## ORIGIN

Query Match 77.1%; Score 16.2; DB 12; Length 328;  
Best Local Similarity 85.7%; Pred. No. 2.3e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 TCGTGAACGTTGAGATGAT 21  
25 TCGTGAACGTTGAGATGAT 45

## Db

RESULT 32  
CD177754/C 329 bp mRNA linear EST 14-SEP-2003  
LOCUS CD177754.1  
DEFINITION MS1-0011T-D106-F07-U.G MS1-001 Schistosoma mansoni cDNA clone  
MS1-0011T-D106-F07.G, mRNA sequence.  
ACCESSION CD177754  
VERSION CD177754.1 GI:34708220  
KEYWORDS EST.  
SOURCE Schistosoma mansoni  
ORGANISM Schistosoma mansoni

REFERENCE Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae; Schistosoma.  
1 (bases 1 to 329)  
AUTHORS Verjovski-Almeida, S., Demarco, R., Martins, E.A.L., Guimaraes, P.E.M., Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y., Jr., Kitajima, J.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F., Coutson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L., Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A., Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A., Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T., Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M., Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.  
TITLE Transcription analysis of the acclomate human parasite Schistosoma mansoni  
JOURNAL Nat. Genet. 35 (2), 148-157 (2003)  
MEDLINE 22879926  
COMMENT Contact: Dr. Sergio Verjovski-Almeida  
Departamento de Bioquímica  
Instituto de Química - Universidade de São Paulo  
Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 São Paulo - SP, Brasil  
Tel: +55-11-3091-2173  
Fax: +55-11-3091-2166  
Email: verjovski@usp.br  
This sequence was derived from the PAPESP Schistosoma mansoni EST Genome Project. All sequences in the project were assembled and annotated. This entry and all the assembled sequences can be seen in the following URL: <http://bioinfo.iq.usp.br/schisto/>  
Plate: MS1-0011T-D106 row: 7 column: F.

## FEATURES

source  
1..329  
location/Qualifiers  
/organism="Schistosoma mansoni"  
/mol\_type="mRNA"  
/db\_xref="taxon:6183"  
/clone="MS1-0011T-D106-F07.G"  
/sex="mixed pool"  
/dev\_stage="schistosomulum"  
/lab\_host="in vitro culture"  
/clone\_lib="MS1-0011"  
/note="Vector: pGEM T-easy"

## ORIGIN

Query Match 77.1%; Score 16.2; DB 14; Length 329;  
Best Local Similarity 85.7%; Pred. No. 2.3e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 TCGTGAACGTTGAGATGAT 21  
278 TCGTGAACGTTGAGATGAT 258

## Db

RESULT 33  
CD092357/C 342 bp mRNA linear EST 14-SEP-2003  
LOCUS CD092357.1  
DEFINITION MC1-0096T-D093-B07-U.G MC1-0096 Schistosoma mansoni cDNA clone  
MC1-0096T-D093-B07.G, mRNA sequence.  
ACCESSION CD092357  
VERSION CD092357.1 GI:34643083  
KEYWORDS EST.  
SOURCE Schistosoma mansoni  
ORGANISM Schistosoma mansoni

REFERENCE Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae; Schistosoma.  
1 (bases 1 to 342)  
AUTHORS Verjovski-Almeida, S., Demarco, R., Martins, E.A.L., Guimaraes, P.E.M., Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y., Jr., Kitajima, J.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F., Coutson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L., Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A., Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A., Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T., Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M., Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.  
TITLE Transcription analysis of the acclomate human parasite Schistosoma mansoni  
JOURNAL Nat. Genet. 35 (2), 148-157 (2003)  
MEDLINE 22879926  
COMMENT Contact: Dr. Sergio Verjovski-Almeida  
Departamento de Bioquímica  
Instituto de Química - Universidade de São Paulo  
Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 São Paulo - SP, Brasil  
Tel: +55-11-3091-2173  
Fax: +55-11-3091-2166  
Email: verjovski@usp.br  
This sequence was derived from the PAPESP Schistosoma mansoni EST Genome Project. All sequences in the project were assembled and annotated. This entry and all the assembled sequences can be seen in the following URL: <http://bioinfo.iq.usp.br/schisto/>  
Plate: MC1-0096T-D093 row: 7 column: B.

## FEATURES

source  
1..342  
location/Qualifiers  
/organism="Schistosoma mansoni"  
/mol\_type="mRNA"  
/db\_xref="taxon:6183"  
/clone="MC1-0096T-D093-B07.G"  
/sex="mixed pool"  
/dev\_stage="cercaria"  
/lab\_host="Biomphalaria glabrata"  
/clone\_lib="MC1-0096"  
/note="Vector: pGEM T-easy"

## ORIGIN

Query Match 77.1%; Score 16.2; DB 14; Length 342;  
Best Local Similarity 85.7%; Pred. No. 2.3e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 TCGTGAACGTTGAGATGAT 21  
131 TCGTGAACGTTGAGATGAT 111

## Db

RESULT 34  
A0840347 374 bp DNA linear GSS 01-SEP-1999

## ORIGIN

**DEFINITION** nbxb0055D20f CUGI Rice BAC Library Oryza sativa (japonica cultivar-group) genomic clone nbxb0055D20f, genomic survey sequence.

**ACCESSION** AQ840347

**VERSION** AQ840347.1 GI:58184400

**KEYWORDS** GSS.

**SOURCE** Oryza sativa (japonica cultivar-group)

**ORGANISM** Oryza sativa (japonica cultivar-group)

**REFERENCE** Eukaryota, Viridiplantae, Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Ehrhartoideae; Oryzaceae; Oryza.

**AUTHORS** Wing, R.A. and Dean, R.A.

**TITLE** A BAC End Sequencing Framework to Sequence the Rice Genome

**JOURNAL** Unpublished (1998)

**COMMENT** Contact: Wing RA  
Clemson University Genomics Institute  
Clemson University  
100 Jordan Hall, Clemson, SC 29634, USA  
Tel: 864 656 7288  
Fax: 864 656 4293  
Email: rwing@clemson.edu  
Seq primer: TAATACGACTCACTATAGG3  
Class: BAC ends  
High quality sequence stop: 314.

**FEATURES**

**source**

1..374  
location/Qualifiers

/organism="Oryza sativa (japonica cultivar-group)"  
/mol\_type="genomic DNA"  
/strain="Japonica"  
/cultivar="Nipponbare"  
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/clone="nbxb0055D20f"  
/issue\_type="leaf"  
/lab\_host="E. coli DH10B"  
/clone\_1fb="CUGI Rice BAC Library"  
/note="Vector: pBELOBAC11; Site 1: HindIII; Site 2: HindIII; Rice is one of two most popular grains in the world. Half of the world population especially those inhabiting highly populated areas of the humid tropics and subtropics, rely on rice as their primary source of carbohydrate. Monocotyledonous rice is a diploid plant (2n=24) with a haploid genome equivalent of 431 Mbp (Arumuganathan and Earle, 1991). The relatively small genome of rice, three times larger than that of Arabidopsis, makes it suitable for genomic studies. In order to facilitate positional cloning, physical mapping and genome sequencing of rice, we have constructed a BAC library from Oryza sativa, Nipponbare variety. The library contains 36,864 clones with an average insert size of 128.5 Kb providing 10.9 haploid genome equivalents. The deep coverage allows the isolation a particular sequence with a probability of 99.9%. Two high density filters, each containing 18,432 clones (doubly spotted), represent the whole library for colony screening."

**ORIGIN**

Query Match 77.1%; Score 16.2; DB 28; Length 374;  
Best Local Similarity 85.7%; Pred. No. 2.4e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

**QY**

1 TCGTCGACGCTTCGATGAT 21  
180 TTGTCACGCTTCGATGAT 200

**Db**

CD190376 375 bp mRNA linear EST 14-SEP-2003  
LOCUS CD190376/c  
DEFINITION MS1-0064U-A245-H12-U-B MS1-0064 Schistosoma mansoni cDNA clone  
ACCESSION CD190376  
VERSION CD190376.1 GI:34720349

**KEYWORDS** EST.

**SOURCE** Schistosoma mansoni

**ORGANISM** Schistosoma mansoni

**REFERENCE** Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea; Strigeiida; Schistosomatidae; Schistosomidae; Schistosoma.

**AUTHORS** 1 (bases 1 to 375)  
Verjovski-Almeida, S., Demarco, R., Martins, E.A.L., Guimaraes, P.E.M., Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y., Jr., Kitajima, J.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F., Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L., Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A., Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A., Sa, R.G., Stuckert, G.C., Soares, M.B., Gargioli, C., Kawano, T., Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Wenck, C.F.M., Senubal, J.C., Leite, L.C.C. and Dias-Neto, E.

**TITLE** Transcriptional analysis of the acelomate human parasite Schistosoma mansoni

**JOURNAL** Nat. Genet. 35 (2), 148-157 (2003)

**MEDLINE** 22879926

**COMMENT** Contact: Dr. Sergio Verjovski-Almeida  
Departamento de Biogenética  
Instituto de Química - Universidade de São Paulo  
Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 São Paulo - SP, Brasil  
Tel: +55-11-3091-2173  
Fax: +55-11-3091-2186  
Email: verjovski@usp.br

This sequence was derived from the FAPESP Schistosoma mansoni EST Genome Project. All sequences in the project were assembled and annotated. This entry and all the assembled sequences can be seen in the following URL <http://bioinfo.iq.usp.br/schisto/>  
Plate: MS1-0064U-A245 row: 12 column: H.

**FEATURES**

**source**

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location/Qualifiers

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/dev\_stage="schistosomulum"  
/lab\_host="in vitro culture"  
/clone\_1fb="MS1-0064"  
/note="Vector: pGEM T-easy"

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**Db**

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VERSION CD178552.1 GI:34709002

**QY**

1 TCGTCGACGCTTCGATGAT 21  
116 TCGTCTACTTCGACATGAT 96

**Db**

CD178552 377 bp mRNA linear EST 14-SEP-2003  
LOCUS CD178552/c  
DEFINITION MS1-0014P-A231-C01-U-B MS1-0014 Schistosoma mansoni cDNA clone  
ACCESSION CD178552  
VERSION CD178552.1 GI:34709002

**QY**

1 TCGTCGACGCTTCGATGAT 21  
116 TCGTCTACTTCGACATGAT 96

**Db**

CD178552 377 bp mRNA linear EST 14-SEP-2003  
LOCUS CD178552/c  
DEFINITION MS1-0014P-A231-C01-U-B MS1-0014 Schistosoma mansoni cDNA clone  
ACCESSION CD178552  
VERSION CD178552.1 GI:34709002

**QY**

1 TCGTCGACGCTTCGATGAT 21  
116 TCGTCTACTTCGACATGAT 96

**Db**

CD178552 377 bp mRNA linear EST 14-SEP-2003  
LOCUS CD178552/c  
DEFINITION MS1-0014P-A231-C01-U-B MS1-0014 Schistosoma mansoni cDNA clone  
ACCESSION CD178552  
VERSION CD178552.1 GI:34709002

**QY**

1 TCGTCGACGCTTCGATGAT 21  
116 TCGTCTACTTCGACATGAT 96

**Db**

CD178552 377 bp mRNA linear EST 14-SEP-2003  
LOCUS CD178552/c  
DEFINITION MS1-0014P-A231-C01-U-B MS1-0014 Schistosoma mansoni cDNA clone  
ACCESSION CD

Sa.R.G., Stukart,G.C., Soares,M.B., Gargioni,C., Kawano,T.,  
Rodrigues,V., Madeira,A.M.B.N., Wilson,R.A., Menck,C.F.M.,  
Setubal,J.C., Leite,L.C.C. and Dias-Neto,E.  
Transcriptome analysis of the acelomate human parasite Schistosoma  
mansoni

TITLE  
JOURNAL  
MEDLINE  
COMMENT  
Nat. Genet. 35 (2), 148-157 (2003)

CONTACT: Dr. Sergio Verjovski-Almeida  
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Fax: +55-11-3091-2186  
Email: verjovsk@usp.br  
This sequence was derived from the FAPESP Schistosoma mansoni EST  
Genome Project. All sequences in the project were assembled and  
annotated. This entry and all the assembled sequences can be seen  
in the following URL: <http://bioinfo.iq.usp.br/schisto/>  
Plate: MSI-0014P-A231 row: 1 column: C.

FEATURES  
source

1.377  
Location/Qualifiers

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/dev\_stage="schistosomulum"  
/lab\_host="in vitro culture"  
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ORIGIN

Query Match 77.1%; Score 16.2; DB 14; Length 377;  
Best Local Similarity 85.7%; Pred. No. 2.4e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TCGTGAACGTTGCGATGAT 21  
DB 140 TCAATCAACGGTCGATGAT 120

RESULT 37  
LOCUS CD187006/c 384 bp mRNA linear EST 14-SEP-2003  
DEFINITION MSI-0056U-V332-D10-U.B MSI-0056 Schistosoma mansoni cDNA clone  
MSI-0056U-V332-D10.B, mRNA sequence.

ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
EST

Schistosoma mansoni  
Schistosoma mansoni  
Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;  
Strigeidae; Schistosomatidae; Schistosomatidae; Schistosoma.  
1 (bases 1 to 384)

REFERENCE  
AUTHORS  
Verjovski-Almeida,S., Demarco,R., Martins,E.A.L., Guimaraes,P.E.M.,  
Ojopi,B.P.B., Paquola,A.C.M., Piazza,J.P., Nishiyama,M.Y. Jr.,  
Kitajima,J.P., Adamson,R.E., Ashton,P.D., Bonaldo,M.F.,  
Coulson,P.S., Dillon,G.P., Farias,L.P., Gregorio,S.P., Ho,P.L.,  
Leite,R.A., Malaguas,L.C.C., Marques,R.C.P., Miyasato,P.A.,  
Nascimento,A.L.T.O., Ohlweiler,F.P., Reis,E.M., Ribeiro,M.A.,  
Sa,R.G., Stukart,G.C., Soares,M.B., Gargioni,C., Kawano,T.,  
Rodrigues,V., Madeira,A.M.B.N., Wilson,R.A., Menck,C.F.M.,  
Setubal,J.C., Leite,L.C.C. and Dias-Neto,E.  
Transcriptome analysis of the acelomate human parasite Schistosoma  
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TITLE  
JOURNAL  
MEDLINE  
COMMENT  
Nat. Genet. 35 (2), 148-157 (2003)

CONTACT: Dr. Sergio Verjovski-Almeida  
Departamento de Biogenética - Universidade de São Paulo  
Instituto de Química - Universidade de São Paulo, 05508-900 São Paulo - SP,  
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Email: verjovsk@usp.br  
This sequence was derived from the FAPESP Schistosoma mansoni EST  
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in the following URL: <http://bioinfo.iq.usp.br/schisto/>  
Plate: MSI-0056U-V332 row: 10 column: D.

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/sex="mixed pool"  
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/note="Vector: pCEM T-easy"

ORIGIN

Query Match 77.1%; Score 16.2; DB 14; Length 384;  
Best Local Similarity 85.7%; Pred. No. 2.4e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TCGTGAACGTTGCGATGAT 21  
DB 136 TCGTCAACTTCGACATGAT 116

RESULT 38  
LOCUS CD190545/c 384 bp mRNA linear EST 14-SEP-2003  
DEFINITION MSI-0064U-A263-D10-U.B MSI-0064 Schistosoma mansoni cDNA clone  
MSI-0064U-A263-D10.B, mRNA sequence.

ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
EST

Schistosoma mansoni  
Schistosoma mansoni  
Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;  
Strigeidae; Schistosomatidae; Schistosomatidae; Schistosoma.  
1 (bases 1 to 384)

REFERENCE  
AUTHORS  
Verjovski-Almeida,S., Demarco,R., Martins,E.A.L., Guimaraes,P.E.M.,  
Ojopi,B.P.B., Paquola,A.C.M., Piazza,J.P., Nishiyama,M.Y. Jr.,  
Kitajima,J.P., Adamson,R.E., Ashton,P.D., Bonaldo,M.F.,  
Coulson,P.S., Dillon,G.P., Farias,L.P., Gregorio,S.P., Ho,P.L.,  
Leite,R.A., Malaguas,L.C.C., Marques,R.C.P., Miyasato,P.A.,  
Nascimento,A.L.T.O., Ohlweiler,F.P., Reis,E.M., Ribeiro,M.A.,  
Sa,R.G., Stukart,G.C., Soares,M.B., Gargioni,C., Kawano,T.,  
Rodrigues,V., Madeira,A.M.B.N., Wilson,R.A., Menck,C.F.M.,  
Setubal,J.C., Leite,L.C.C. and Dias-Neto,E.  
Transcriptome analysis of the acelomate human parasite Schistosoma  
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TITLE  
JOURNAL  
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COMMENT  
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Tel: +55-11-3091-2173  
Fax: +55-11-3091-2186  
Email: verjovsk@usp.br  
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in the following URL: <http://bioinfo.iq.usp.br/schisto/>  
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FEATURES  
source

1.384  
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/clone="MS1-0064U-A263-D10.B"  
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 /lab\_host="in vitro culture"  
 /clone\_lib="MS1-0064"  
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## ORIGIN

Query Match 77.1%; Score 16.2; DB 14; Length 384;  
 Best Local Similarity 85.7%; Pred. No. 2.4e+03;  
 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 TCGTGAACGTCGAGATGAT 21  
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 Db 135 TCGTGAACGTCGAGATGAT 115

RESULT 39  
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 DEFINITION ME1-0084T-L228-D07-U-B ME1-0084 Schistosoma mansoni cDNA clone  
 ACCESSION ME1-0084T-L228-D07.B. mRNA sequence.  
 VERSION CD124011  
 KEYWORDS CD124011.1 GI:34662045  
 SOURCE EST.  
 ORGANISM Schistosoma mansoni  
 Schistosoma mansoni  
 Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;  
 Strigylidae; Schistosomatidae; Schistosomidae; Schistosoma.  
 1 (bases 1 to 392)

REFERENCE  
 AUTHORS Verjovski-Almeida, S., Demarco, R., Martins, E.A.L., Guimaraes, P.E.M.,  
 Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y. Jr.,  
 Kitejima, U.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F.,  
 Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L.,  
 Leite, R.A., Malaguides, L.C.C., Marques, R.C.P., Miyasato, P.A.,  
 Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A.,  
 Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T.,  
 Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M.,  
 Seidubai, J.C., Leite, L.C.C. and Dias-Neto, E.  
 Transcription analysis of the acclimated human parasite Schistosoma  
 mansoni  
 Nat. Genet. 35 (2), 148-157 (2003)

## TITLE

JOURNAL  
 MEDLINE  
 COMMENT 22879926  
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 Brasil  
 Tel: +55-11-3091-2173  
 Fax: +55-11-3091-2186  
 Email: verjovski@iq.usp.br

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 in the following URL: <http://biinfo.iq.usp.br/schisto/>  
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 Location/Qualifiers

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 /organism="Schistosoma mansoni"  
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 /note="Vector: pGEM T-easy"

## ORIGIN

Query Match 77.1%; Score 16.2; DB 14; Length 392;  
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 Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 TCGTGAACGTCGAGATGAT 21  
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 Db 278 TCGTGAACGTCGAGATGAT 258

RESULT 40  
 AM699669/c 410 bp mRNA linear EST 18-APR-2000  
 DEFINITION gb30d06.y1 Moss EST library PPN Physcomitrella patens cDNA clone  
 PEP SOURCE ID:PPN140212.5 similar to gb:emb|22315.1|ATBPS18A  
 A.thaliana ribosomal protein gene (PLANT); mRNA sequence.  
 AM699669  
 1 (bases 1 to 410)

ACCESSION AM699669.1 GI:7583764  
 VERSION EST.  
 KEYWORDS Physcomitrella patens  
 SOURCE Physcomitrella patens  
 ORGANISM Physcomitrella patens  
 Bryopsida; Viridiplantae; Streptophyta; Embryophyta; Bryophyta;  
 Eukaryota; Viridiplantae; Funariidae; Funariaceae; Physcomitrella.  
 1 (bases 1 to 410)  
 Quatrano, R., Bashardes, S., Cove, D., Cuning, A., Knight, C.,  
 Clifton, S., Marra, M., Hillier, L., Pape, D., Martin, J., Wylie, T.,  
 Underwood, K., Theising, B., Allen, M., Bowers, Y., Person, B.,  
 Swaller, T., Steptoe, M., Gibbons, M., Harvey, N., Ritter, E.,  
 Jackson, Y., McCann, R., Waterson, R. and Wilson, R.  
 Leeds/Wash U Moss EST Project  
 Unpublished (1999)

## TITLE

JOURNAL  
 COMMENT  
 Contact: Ralph Quatrano  
 Leeds/Wash U Moss EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: est@watson.wustl.edu

The libraries were constructed by Dr. Stevros Bashardes as part of the  
 Physcomitrella EST program (PEP) at the Univ. of Leeds (UK) and  
 Washington Univ. in St. Louis (USA) DNA sequencing by Washington  
 University Genome Sequencing Center For information on obtaining a  
 clone please contact: Celia Knight (c.d.knight@leeds.ac.uk)  
 Seq primer: -40RP from Gibco.  
 Location/Qualifiers

## FEATURES

source

1.410  
 /organism="Physcomitrella patens"  
 /mol\_type="mRNA"  
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 /lab\_host="DH10B"  
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 XhoI. Construction of the cDNA library was carried out  
 using Stratagene's 'unizap - cDNA synthesis kit'. cDNA was  
 constructed using an oligo dT primer/linker that contains  
 a XhoI site within it. Following ds cDNA synthesis,  
 EcoRI adaptors were ligated to the blunt ends and sample  
 was digested with XhoI. The result is cDNA with an EcoRI  
 sticky end on one side and a XhoI sticky end on the other.  
 This cDNA was ligated directionally in unizap arms. The  
 vector is designed containing the pBluescript sequence as  
 well as lambda DNA and cDNA is cloned within this  
 pBluescript sequence. The vector was then packaged using  
 Gold gigacloning extracts. Library was grown in XlBlue  
 MRF cells and amplified. The library was excised by mass  
 excision using Stratagene 'mass excision kit' that uses  
 exsist as a helper phage that releases the pBluescript  
 sequence and circularises it as single stranded plasmids  
 that are then packaged (by helper phage) and secreted out  
 of the host cell as phagemids. SOLR cells were transformed  
 with phagemids and the library was plated out on LB-amp  
 plates to select for transformants. Approximately  
 1,000,000 colonies were grown and recovered. The double  
 stranded plasmid library was recovered by using Qulagen  
 Midi prep kit. 2 micro grams of each library were used to  
 transform DH10B cells by electroporation."



## ORIGIN

Query Match	77.1%;	Score 16.2;	DB 10;	Length 410;
Best Local Similarity	85.7%;	Pred. No. 2.5e+03;		
Matches 18;	Conservative 0;	Mismatches 3;	Indels 0;	Gaps 0;

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Qy      1 TCGTGACGTCGAGATGAT 21
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Db      87 TCTTCAACGTCGAGATCAT 67

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Job time : 3213 secs

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